

Application
for
United States Letters Patent

To all whom it may concern:

Be it known that we, Moses Rodriguez and Daren Ure

have invented certain new and useful improvements in

TREATMENT OF CENTRAL NERVOUS SYSTEM DISEASES BY ANTIBODIES AGAINST GLATIRAMER ACETATE

of which the following is a full, clear and exact description.

**TREATMENT OF CENTRAL NERVOUS SYSTEM
DISEASES BY ANTIBODIES AGAINST GLATIRAMER ACETATE**

This application claims the benefit of U.S. Provisional No. 60/287,171, filed April 27, 2001, U.S. Provisional No. 60/269,788, filed February 16, 2001, and U.S. Provisional No. 60/212,577, filed June 20, 2000, the contents of which are hereby incorporated by reference into the present application.

Throughout this application, various references are referenced by arabic numbers within parenthesis. Full citations for these references may be found at the end of the specification, immediately preceding the claims. These references, in their entireties, are hereby incorporated by reference to more fully describe the state of the art to which this invention pertains.

Field of the Invention

The present invention is directed to the treatment of central nervous system (CNS) diseases by antibodies.

Background of the Invention

The nervous system of vertebrates is divided into the central nervous system, comprised of the brain and spinal cord, and the peripheral nervous system, consisting of the outlying nerves (16). The axons of most nerve cells are covered with a myelin sheath, a stack of specialized plasma membranes. Glial cells that wrap around the axons produce the myelin sheath. In the CNS, these cells are called oligodendrocytes. The myelin membranes of the CNS contain myelin basic protein (MBP) and a proteolipid (PLP) that is not found elsewhere in vertebrates. Each region of myelin formed by an individual glial cell is separated from the next region by an

unmyelinated area called the node of Ranvier; only at nodes is the axonal membrane in direct contact with the extracellular fluid.

5 The myelin sheath, which can be 10-12 myelin wraps thick, acts as an electric insulator of the axon by preventing the transfer of ions between the axonal cytoplasm and the extracellular fluids (16). Thus all electric activity in axons is confined to the nodes of Ranvier, the sites where 10 ions can flow across the axonal membrane. Node regions contain a high density of voltage-dependent Na^+ channels, about 10,000 per μm^2 , whereas the regions of axonal membrane between the nodes have few if any channels.

15 The excess cytosolic positive ions generated at a node during the membrane depolarization associated with an action potential diffuse through the axonal cytoplasm to the next node with very little loss or attenuation because ions are capable of moving across the axonal membrane only at the 20 myelin-free nodes (16). Thus a depolarization at one node spreads rapidly to the next node, and the action potential jumps from node to node. For this reason, the conduction velocity of myelinated nerves is much greater than that of 25 unmyelinated nerves of the same diameter. For example, a 12- μm -diameter myelinated vertebrate axon and a 600- μm -diameter unmyelinated squid axon both conduct impulses at 12m/s.

One of the more common neurologic diseases in human adults is 30 multiple sclerosis. This condition is a chronic, frequently progressive, inflammatory CNS disease characterized pathologically by primary demyelination. The etiology and pathogenesis of multiple sclerosis are unknown. Researchers have hypothesized that multiple sclerosis is an autoimmune disease (14, 23, 47) or that a virus, bacteria or other 35 agent, precipitates an inflammatory response in the CNS,

which leads to either direct or indirect ("bystander") myelin destruction, potentially with an induced autoimmune component (31, 38). Thus, a rebuilding of the myelin sheath, or remyelination, can treat multiple sclerosis.

5

Spontaneous remyelination of axons within lesions by oligodendrocytes has been shown to occur to a small degree in SJL/J mice and multiple sclerosis patients (1). Several types of antibodies have been found to promote remyelination 10 (1). Some of these antibodies are polyclonal, derived by immunization with spinal cord homogenate or myelin basic protein (71). One remyelination-promoting antibody is monoclonal (SCH 94.03) (1). The isotype of these antibodies is IgM, and they share the characteristic of binding to the 15 surface of oligodendrocytes (1). Also, they are polyreactive, binding to a variety of cytoskeletal proteins or proteins with repeating structures (1).

Of clinical importance is the question whether morphologic 20 regeneration of thin myelin sheaths contributes to functional recovery (1). Computer simulations indicate that new myelin formation even by inappropriately thin sheaths improves impulse conduction (1). Since the axon membrane of normally myelinated fibers is highly differentiated, it is necessary 25 for sodium channels to be present at high density at the node of Ranvier to propagate saltatory conduction. Experimental evidence suggests that newly formed nodes do develop the required high sodium channel density as demonstrated by saxitoxin binding. Data suggest that remyelination even by 30 inappropriately thin myelin improves conduction in a previously demyelinated axon. Therefore, any strategy to promote this morphologic phenomenon has the potential of producing functional recovery. Studies examining biopsy tissues from patients with severe acute exacerbations 35 demonstrate that demyelination is a significant component of

the acute multiple sclerosis lesion (57). Therefore, remissions are probably associated with significant CNS remyelination (1).

5 One commonly utilized experimental model of multiple sclerosis is induced by Theiler's murine encephalomyelitis virus (TMEV) (15, 59). In the TMEV model, spinal cord demyelination is influenced by the immune response to virus infection and is therefore continuously sensitive to
10 immunomodulation. Previous experiments in Strain Jackson Laboratories (SJL) mice infected with TMEV showed that 4 to 5% of the demyelinated area exhibited significant spontaneous remyelination (62). In protocols using antibody therapy and monoclonal antibody therapy, this number increased up to 30-
15 35% (41, 58, 71). For instance, using the TMEV model, it was demonstrated that the passive transfer of CNS specific antiserum (63) and purified antibodies (55, 62, 71) directed against myelin components promoted CNS remyelination. This contrasts with the conventional view that the humoral immune
20 response plays a pathogenic role in CNS demyelination (56). Researchers also generated a monoclonal antibody that reacted against a surface component of oligodendrocytes and promoted remyelination (40-42). It has also been shown that antibodies reactive with myelin basic protein (MBP) promoted
25 CNS remyelination (58). In these experiments, infected SJL mice were treated with the whole anti-serum or affinity purified mouse antibodies directed against rabbit or rat myelin basic proteins. There was extensive evidence for new myelin synthesis as measured by quantitative morphometry.
30 Electron microscopy revealed numerous oligodendrocytes and remyelinated CNS axons with a relative lack of inflammatory cells. Viral antigen persisted in these animals despite enhanced CNS remyelination. These findings indicated for the first time that antibodies reactive against a myelin
35 autoantigen and in particular, MBP, have the potential for

myelin repair.

U.S. Patent 5,591,629 describes the promotion of CNS remyelination in the TMEV model through SCH 94.03 monoclonal antibodies directed against spinal cord homogenate (SCH) (1). SCH encompasses myelin antigens, such as MBP (64) and proteolipid protein (PLP) (12, 67). Although SCH contains MBP, this antibody does not react with MBP. The SCH 94.03 antibody is an IgM which recognizes cytoplasmic determinants on glial cells. It also recognizes surface determinants on glial cells, including oligodendrocytes. Experiments demonstrated that the antibody does not react with TMEV. In addition, the antibody was shown to promote the proliferation of glial cells in mixed rat brain culture in a dose-dependent manner. SCH 94.03 is a natural autoantibody.

A treatment that has been shown to be effective in reducing exacerbations of multiple sclerosis is the administration of glatiramer acetate (2-6, 31). Daily subcutaneous injections of glatiramer acetate (20 mg/injection) reduce relapse rates, appearance of new lesions by magnetic resonance imaging (MRI), and progression of disability (26). COPAXONE® is the brand name for glatiramer acetate (also known as Copolymer-1 (77), Copolymer 1, Cop-1 or Cop), an FDA-approved drug for the treatment of multiple sclerosis. Glatiramer acetate, the active ingredient of COPAXONE®, consists of the acetate salts of synthetic polypeptides, containing four naturally occurring amino acids: L-glutamic acid, L-alanine, L-tyrosine, and L-lysine (77) with an average molar fraction of L-glutamic acid: 0.129-0.153; L-alanine: 0.392-0.462; L-tyrosine: 0.086-0.100; L-lysine: 0.300-0.376, respectively. The average molecular weight of glatiramer acetate is 4,700-11,000 daltons (77). Chemically, glatiramer acetate is designated L-glutamic acid polymer with L-alanine, L-lysine and L-tyrosine, acetate (salt) (77). Its structural formula

is:

$$(\text{Glu, Ala, Lys, Tyr})_x \cdot \text{CH}_3\text{COOH}$$

$$(C_5H_9NO_4 \cdot C_3H_7NO_2 \cdot C_6H_{14}N_2O_2 \cdot C_9H_{11}NO_3)_x \cdot xC_2H_4O_2$$

CAS - 147245-92-9

(77). Glatiramer acetate is also written as: poly[L-Glu¹³⁻¹⁵, L-Ala³⁹⁻⁴⁶, L-Tyr⁸⁻⁶⁻¹⁰, L-Lys³⁰⁻³⁷] nCH₃COOH.

10 Unlike myelin basic protein (MBP), with which it shares some structural characteristics, glatiramer acetate inhibits rather than induces experimental autoimmune encephalomyelitis (EAE), an animal model of MS (37, 65-66). Glatiramer acetate-reactive, type 2 helper T lymphocytes confer
15 resistance to EAE.

In spite of the experimental evidence that glatiramer acetate down-regulates certain immune functions, clinical use of glatiramer acetate indicates that other immune functions are stimulated by the peptide treatment. In rodents, monoclonal antibodies to glatiramer acetate have been generated, some of which cross-react with MBP (68), but other cross-reactivities are unknown. The humoral response to glatiramer acetate may have diverse roles in multiple sclerosis. Some autoreactive antibodies to myelin antigens might contribute to pathogenesis (22, 35). Other antibodies, such as those that develop in a subset of interferon-treated patients, may neutralize therapeutic efficacy. A third possibility is that some antibodies may in fact be protective. All individuals have antibodies to a wide range of endogenous antigens, including MBP, suggesting that natural autoantibodies represent a conserved adaptation to nervous system disease and trauma. In support of a protective role for autoreactive antibodies, mouse or human antibodies reactive to the central nervous system (CNS) have been found to promote myelin repair

in viral experimental model of multiple sclerosis (41, 48, 53, 58). Antibodies have been found to stimulate remyelination in SJL mice that were chronically infected with EAE (41, 58).

5 Antisera against glatiramer acetate have been employed to
investigate the mechanism by which L-glatiramer acetate is
effective against Experimental Allergic Encephalomyelitis
(EAE) (74-75). For this purpose, Webb et al. measured the
cross-reactivity of L-glatiramer acetate anti-sera with D-
10 glatiramer acetate and Copolymer 4 (L-glatiramer acetate
modified by the replacement of tyrosine with tryptophan)
(75). Webb et al. carried out a similar experiment to
determine the reactivity of L-glatiramer acetate anti-sera
with L-glatiramer acetate, and the cross-reactivity of L-
15 glatiramer acetate anti-sera with AGT (alanine, glutamic acid
and tyrosine), BE (Basic Encephalitogen), AAspLT (alanine,
aspartic acid, lysine and tyrosine) and AGL (alanine,
glutamic acid and lysine) (74).

20 Monoclonal antibodies against glatiramer acetate and against
MBP have also been utilized to probe the mechanism of
glatiramer acetate in treatment of EAE (68). The cross-
reactivity of monoclonal antibodies against glatiramer
acetate with MBP was analyzed by Teitelbaum et al (68). They
25 also determined the cross-reactivity of monoclonal antibodies
against MBP with glatiramer acetate (68). Another focus of
their experiments was the cross-reactivity of glatiramer
acetate anti-sera with MBP and of MBP-antisera with
glatiramer acetate (68). The cross-reactivity of anti-MBP
30 anti-sera with glatiramer acetate was additionally
investigated by Lisak et al (37).

Summary of the Invention

The subject invention concerns a humanized antibody directed against an epitope on glatiramer acetate, also known as Copolymer 1, Copolymer-1, Cop-1 or Cop.

5

The subject invention further encompasses a F_{ab} fragment that binds to an epitope on glatiramer acetate.

10 In addition, the subject invention relates to a pharmaceutical composition comprising an antibody directed against an epitope on glatiramer acetate in an amount effective to treat a central nervous system disease and a pharmaceutically acceptable carrier.

15 The subject invention also provides a method of stimulating remyelination of central nervous system axons comprising contacting the axons with an antibody directed against an epitope on glatiramer acetate in an amount effective to stimulate remyelination of central nervous system axons.

20

The subject invention additionally includes a method of treating a subject suffering from a disease associated with demyelination of central nervous system axons comprising administering to the subject an effective amount of an antibody directed against an epitope on glatiramer acetate in an amount effective to treat the disease associated with demyelination of central nervous system axons.

25 The subject invention further relates to a method of stimulating remyelination of central nervous system axons comprising contacting the axons with glatiramer acetate in an amount effective to stimulate remyelination of central nervous system axons.

30 35 The subject invention also concerns a method of treating a

subject suffering from a disease associated with demyelination of central nervous system axons comprising administering to the subject glatiramer acetate in an amount effective to treat the disease associated with demyelination
5 of central nervous system axons, wherein the disease associated with demyelination of central nervous system axons is selected from the group consisting of: acute disseminated encephalomyelitis, transverse myelitis, demyelinating genetic diseases, spinal cord injury, virus-induced demyelination,
10 Progressive Multifocal Leucoencephalopathy, Human Lymphotrophic T-cell Virus I (HTLV-I)-associated myelopathy, and nutritional metabolic disorders.

Finally, the subject invention encompasses a method of
15 stimulating proliferation of lymphocytes comprising contacting the lymphocytes with an antibody directed against an epitope on glatiramer acetate in an amount effective to stimulate lymphocyte proliferation.

20 **Description of the Drawings**

Figure 1 demonstrates that glatiramer acetate does not alter the extent of spinal cord demyelinating lesions during early disease. Each dot represents one mouse.

Figure 2 shows that glatiramer acetate at high doses increases the extent of spinal cord demyelinating lesions during late disease. Each dot represents one mouse.

Figure 3 reveals that glatiramer acetate does not alter the extent of remyelination during late disease. Each dot represents one mouse.

30 **Figure 4** shows that glatiramer acetate does not alter the extent of brain pathology during late disease. Each dot represents one mouse.

Figure 5 demonstrates that antibodies against epitopes on glatiramer acetate do not alter the extent of spinal cord

demyelinating lesions during late disease. Each dot represents one mouse.

Figure 6 reveals that antibodies against epitopes on glatiramer acetate promote remyelination during late disease.

5 Each dot represents one mouse.

Figure 7 reports that antibodies against glatiramer acetate do not affect demyelination (Figure 7A), but promote remyelination (Figure 7B).

Figure 8 depicts the glatiramer acetate IgG developed by non-
10 infected mice after immunization.

Figure 9 shows that chronically diseased mice develop antibodies against epitopes on glatiramer acetate in response to glatiramer acetate treatment and that levels of these antibodies are increased by co-administration with incomplete
15 Freund's adjuvant (IFA).

Figure 10 demonstrates that the levels of antibodies against epitopes on glatiramer acetate increase as a function of time post-immunization and as a function of dose in chronically diseased mice.

20 **Figure 11** reveals that glatiramer acetate IgG is detectable in serum 10 days after passive transfer to chronically diseased, non-immunized mice. Individual symbols represent means (\pm SEM) from 4-6 mice, except the untreated group (2 mice).

25 **Figure 12** displays the concentration of glatiramer acetate IgG in serum generally rises with the dosage of glatiramer acetate (Figure 12A). Figure 12B portrays the isotypes of purified glatiramer acetate Ig, Normal Ig and pooled mouse serum Ig and shows that IgG1 was the predominant glatiramer acetate isotype.

30 **Figure 13** shows that affinity-purified antibodies against epitopes on glatiramer acetate (IgG and IgM) have high reactivity to Cop-1 by ELISA, whereas Normal antibodies have no reactivity to glatiramer acetate.

Figure 14 depicts the low polyreactivity of antibodies against epitopes on glatiramer acetate and Normal antibodies. In the figure, Hu=human; Ms=mouse; Rb=rabbit; Bov=bovine; BSA=bovine serum albumin.

5 **Figure 15** shows that antibodies against epitopes on glatiramer acetate and Normal antibodies have a wide spectrum of antibody isotypes.

10 **Figures 16-18** reveal that glatiramer acetate antibodies stimulate proliferation of lymph node-derived lymphocytes from mice immunized with myelin peptides but not from non-immunized mice. The quantities shown are per 200 μ l medium per well.

15 **Figures 19-20** demonstrate that antibodies against epitopes on glatiramer acetate bind to microglia and macrophages, but not to oligodendrocytes. Figures 20-A - 20-D show the staining of antibodies against glatiramer acetate, while Figures 20E - 20-F reflect the staining of O4-positive oligodendrocytes.

Figures 21-22 display demyelinating lesions from chronically diseased, phosphate buffered saline (PBS)-treated mice.

20 **Figure 23** shows CNS reactivity of purified glatiramer acetate Ig. (a, b, c) In rat CNS glial cultures glatiramer acetate IgG bound to the surface of round, non-process-bearing cells located on the upper surface of the cultures. Glatiramer acetate IgG-positive cells co-labeled with the microglial marker, *Bandeiraea simplicifolia* isolectin B₄, but did not co-label with oligodendrocyte markers, A2B5 (immature oligodendrocytes) or O1 (mature oligodendrocytes). (d, e) In spinal cord sections from lesioned spinal cord, biotinylated glatiramer acetate IgG bound in the white matter to glial cells, especially their processes, and to the perivasicular infiltrate, but not to myelin or oligodendrocytes. Gray matter and neuronal cell bodies were also glatiramer acetate IgG-positive. Biotinylated pooled mouse IgG did not bind appreciably to sections.

Figure 24 depicts a cross-section of spinal cord white matter stained for myelin showing normal myelin architecture.

Figures 25-26 portray demyelinating lesions from chronically diseased, glatiramer acetate-treated mice (0.1 mg/injection).

5 Lesions show attempts at remyelination.

Figures 27-28 illustrate demyelinating lesions from chronically diseased, glatiramer acetate antibody-treated mice. Lesions show extensive remyelination (outlined), characterized by thin myelin sheaths around axons.

10 **Figure 29** explains the process of purifying antibodies against epitopes on glatiramer acetate.

Figure 30 outlines the steps by which Normal antibodies are purified.

15 **Detailed Description of the Invention**

The phrase, early disease, is defined as the period up to 45 days post-infection and encompasses the encephalitic stage of disease as well as the beginning of the demyelinating phase of disease (begins at approximately Day 21 post-infection).

20 The phrase, late disease, is defined as the period beyond four months of infection.

25 The phrase, low-dose glatiramer acetate, is defined as single or multiple injections of glatiramer acetate at 0.02-0.1 mg/injection. Given the average weight of adult SJL/J mice of 20 g, a 0.1 mg injection is equivalent to 5 mg/kg.

30 The phrase, high-dose glatiramer acetate, is defined as single or multiple injections of glatiramer acetate at 5 mg/injection. Given the average weight of adult SJL/J mice of 20 g, a 5 mg injection is equivalent to 250 mg/kg.

The subject invention provides a humanized antibody directed

against an epitope on glatiramer acetate.

In one embodiment, this humanized antibody is not cross-reactive with MBP.

5

In another embodiment, this humanized antibody consists essentially of IgG1.

10 In a further embodiment, this humanized antibody does not react with mature oligodendrocytes.

In another embodiment, this humanized antibody cross-reacts with SCH.

15 In one embodiment, this humanized antibody primarily reacts with cells exhibiting a macrophage or microglial phenotype.

In yet another embodiment, this humanized antibody is a monoclonal antibody.

20

In a further embodiment, this humanized antibody is a polyclonal antibody.

25 The subject invention further relates to a F_{ab} fragment that binds to an epitope on glatiramer acetate.

30 In addition, the subject invention involves a pharmaceutical composition comprising an antibody directed against an epitope on glatiramer acetate in an amount effective to treat a demyelinating central nervous system disease and a pharmaceutically acceptable carrier.

In the pharmaceutical composition, this antibody may be a humanized antibody.

35

In the pharmaceutical composition, this antibody may be not cross-reactive with MBP.

5 In the pharmaceutical composition, this antibody may consist essentially of IgG1.

In the pharmaceutical composition, this antibody may not react with mature oligodendrocytes.

10 In the pharmaceutical composition, this antibody may cross-react with SCH.

15 In the pharmaceutical composition, this antibody may primarily react with cells exhibiting a macrophage or microglial phenotype.

In the pharmaceutical composition, this antibody may be a monoclonal antibody.

20 In the pharmaceutical composition, this antibody may be a polyclonal antibody.

25 The subject invention also provides a method of stimulating remyelination of central nervous system axons comprising contacting the axons with an antibody directed against an epitope on glatiramer acetate in an amount effective to stimulate remyelination of central nervous system axons.

30 In this method, the antibody may be a humanized antibody.

35 In this method, the antibody may be not cross-reactive with MBP.

In this method, the antibody may consist essentially of IgG1.

In this method, the antibody may not react with mature oligodendrocytes.

In this method, the antibody may cross-react with SCH.

5

In this method, the antibody may primarily react with cells exhibiting a macrophage or microglial phenotype.

In this method, the antibody may be a monoclonal antibody.

10

In this method, the antibody may be a polyclonal antibody.

Additionally, the subject invention concerns a method of treating a subject suffering from a disease associated with demyelination of central nervous system axons comprising administering to the subject an effective amount of an antibody directed against an epitope on glatiramer acetate in an amount effective to treat the disease associated with demyelination of central nervous system axons.

15

In this method, the antibody may be a humanized antibody directed against an epitope on glatiramer acetate.

20

In this method, the antibody may not be cross-reactive with MBP.

In this method, the antibody may consist essentially of IgG1.

25

In this method, the antibody may not react with mature oligodendrocytes.

In this method, the antibody may cross-reacts with SCH.

30

In this method, the antibody may primarily react with cells exhibiting a macrophage or microglial phenotype.

In this method, the antibody may be a monoclonal antibody.

In this method, the antibody may be a polyclonal antibody.

5 In this method, the disease associated with demyelination of central nervous system axons is selected from the group consisting of: multiple sclerosis, acute disseminated encephalomyelitis, transverse myelitis, demyelinating genetic diseases, spinal cord injury, virus-induced demyelination, 10 Progressive Multifocal Leucoencephalopathy, Human Lymphotrophic T-cell Virus I (HTLV-I)-associated myelopathy, and nutritional metabolic disorders.

In one embodiment, the disease associated with demyelination 15 of central nervous system axons is multiple sclerosis.

In another embodiment, the disease associated with demyelination of central nervous system axons is acute disseminated encephalomyelitis.

20 In an additional embodiment, the disease associated with demyelination of central nervous system axons is transverse myelitis.

25 In a further embodiment, the disease associated with demyelination of central nervous system axons is a demyelinating genetic disease.

30 In yet another embodiment, the disease associated with demyelination of central nervous system axons is a spinal cord injury.

35 In a further embodiment, the disease associated with demyelination of central nervous system axons is virus-induced demyelination.

In another embodiment, the disease associated with demyelination of central nervous system axons is Progressive Multifocal Leucoencephalopathy.

5 In an additional embodiment, the disease associated with demyelination of central nervous system axons is HTLV-associated myelopathy.

10 In another embodiment, the disease associated with demyelination of central nervous system axons is a nutritional metabolic disorder.

In one embodiment, the nutritional metabolic disorder is vitamin B₁₂ deficiency.

15 In another embodiment, the nutritional metabolic disorder is central pontine myelinolysis.

20 In one embodiment, the effective amount is an amount from 0.1 mg to 400 mg.

In a preferred embodiment, the effective amount is an amount from 0.1 mg to 250 mg.

25 In a further embodiment, the effective amount is an amount from 0.5 mg to 400 mg.

In another embodiment, the effective amount is an amount from 0.5 mg to 300 mg.

30 In another embodiment, the effective amount is an amount from 0.5 mg to 250 mg.

35 In further embodiment, the effective amount is an amount from 1.0 mg to 250 mg.

In another embodiment, the effective amount is an amount from 2.5 mg to 225 mg.

In yet another embodiment, the effective amount is an amount 5 from 5.0 mg to 200 mg.

In a further embodiment, the effective amount is an amount from 10 mg to 175 mg.

10 In another embodiment, the effective amount is an amount from 25 mg to 150 mg.

In yet another embodiment, the dosage of antibodies against glatiramer acetate is an amount from 50 mg to 125 mg.

15 In a further embodiment, the effective amount is an amount from 75 mg to 100 mg.

20 The subject invention further provides a method of stimulating remyelination of central nervous system axons comprising contacting the axons with glatiramer acetate in an amount effective to stimulate remyelination of central nervous system axons.

25 The subject invention additionally concerns a method of treating a subject suffering from a disease associated with demyelination of central nervous system axons comprising administering to the subject glatiramer acetate in an amount effective to treat the disease associated with demyelination 30 of central nervous system axons, wherein the disease associated with demyelination of central nervous system axons is selected from the group consisting of: acute disseminated encephalomyelitis, transverse myelitis, demyelinating genetic diseases, spinal cord injury, virus-induced demyelination, 35 Progressive Multifocal Leucoencephalopathy, HTLV-1-associated

myelopathy, and nutritional metabolic disorders.

The subject invention also contains a method of stimulating proliferation of lymphocytes comprising contacting the 5 lymphocytes with an antibody directed against an epitope on glatiramer acetate in an amount effective to stimulate lymphocyte proliferation.

In this method, the antibody may be a humanized antibody 10 directed against an epitope on glatiramer acetate.

In this method, the antibody may be not cross-reactive with MBP.

15 In this method, the antibody may consist essentially of IgG1.

In this method, the antibody may not react with mature oligodendrocytes.

20 In this method, the antibody may cross-react with SCH.

In this method, the antibody may primarily react with cells exhibiting a macrophage or microglial phenotype.

25 In this method, the antibody may be a monoclonal antibody.

In this method, the antibody may be a humanized polyclonal antibody directed against an epitope on glatiramer acetate.

30 In one embodiment, antibodies against glatiramer acetate are generated by intraperitoneal injection of glatiramer acetate into SJL mice. Alternatively, glatiramer acetate could be injected intradermally or intravenously. Other sources of antibodies against glatiramer acetate are contemplated by the 35 invention. These sources include, but are not limited to,

other mice, rabbits, cats, goats, monkeys and humans.

Additionally contemplated by the present invention is a monoclonal antibody directed against an epitope on glatiramer acetate. These antibodies can be created by procedures known to those of skill in the art. Such procedures include, but are not limited to, the creation of hybridomas and antibody libraries.

10 A hybridoma is produced by the fusion of Normal B lymphocytes, which will not grow indefinitely in culture, and myeloma cells, which are immortal (16, 78). The selective medium most often used to culture such fused cells is called HAT medium, because it contains hypoxanthine, aminopterin, 15 and thymidine (16). Normal B lymphocytes can grow in HAT medium, salvage mutants cannot, but their hybrids with Normal B lymphocytes can (16). Mutant myeloma cell lines that have lost the salvage pathways for purines (indicated by their inability to grow in HAT medium) are selected (16). These 20 myeloma cells are then fused with normal B lymphocytes, creating hybridoma cells (16). Like myeloma cells, hybridoma cells can grow indefinitely in culture; like normal B lymphocytes, the fused cells have purine salvage-pathway enzymes and can grow in HAT medium (16). If a mixture of 25 fused and unfused cells is placed in HAT medium, the unfused mutant myeloma cells and the unfused lymphocytes die, leaving a culture of immortal hybridoma cells, each of which produces a monoclonal antibody (16). Clones of hybridoma cells can be tested separately for the production of a desired antibody 30 and the clones containing that antibody then can be cultured in large amounts (16).

Following the cloning of genes encoding antibodies, a library of filamentous phage can be prepared (52). Each phage has 35 the potential to display a unique antibody on its surface,

which is the selectable phenotype (52). Within the phage coat is the genotype that encodes the displayed molecule. This linkage of displayed antibody phenotype with encapsulated genotype via the phage surface forms the basis 5 of the technique (52). Typically, the antibody fragments are displayed on the surface of phage as either F_{ab} fragments, single-chain variable region fragments (scF_{vs}), or dimeric scF_{vs} , also known as diabodies, which differ from scF_{vs} in the reduced length of the linker peptide used and their 10 preference to associate as dimers (52). Library construction is facilitated by the ready availability of phagemid vectors, which allow for construction and display of libraries of these antibody fragments using a single rare cutting restriction enzyme, *SfiI* (52). Selection of antibodies from 15 the library is based on the displayed antibodies' binding specificity and affinity and is generally performed over several rounds of selection and amplification in a process known as panning (52).

20 Phage displayed antibody libraries can be screened by panning on purified antigens immobilized on artificial surfaces or by panning on cell surface expressed antigens (52). In contrast to panning on purified immobilized antigen, cell panning selects for antibodies that are more likely to bind to 25 epitopes *in vivo* (52).

One method of improving the specificity of antibodies in an library is saturated mutagenesis of complementarity determining regions (CDR) (CDR walking mutagenesis) (52). In 30 this approach, saturation mutagenesis of a CDR is constrained to libraries that examine all possible amino acids in the target CDR (52). Two strategies are employed -- either sequential or parallel optimization of CDR (52). In the sequential approach, the library of antibodies with a single 35 randomized CDR is screened by several rounds of panning

against the antigen (52). The selected clone(s) are then used in the construction of a second library where a different single CDR is randomized (52). The panning, selection of clone(s) and construction of a library is 5 repeated several times (52). Sequential optimization takes into account that optimal binding may result from the interdependence of CDRs (52). In the parallel approach, independent libraries are constructed where each library represents the randomization of given CDR (52). Each library 10 is screened by several rounds of panning against the antigen (52). Then, the individually optimized CDRs are combined. If the free energy change of individually optimized CDRs combined is nearly equal to the sum of the free energy changes in the single optimized CDRs, the free energy changes 15 are said to be additive (52). As additivity within the antibody binding site is virtually impossible to predict, Rader and Barbas believe that sequential CDR optimization is preferred over parallel (52).

20 The subject invention further contemplates humanized antibodies against glatiramer acetate. A humanized antibody is a non-human antibody which has been genetically engineered by the substitution of human nucleotide sequences in the nonvariable regions of the non-human antibodies (1, 72). 25 Such substitutions reduce the immunogenicity of the antibodies in humans without significantly lowering the specificity of the antibodies.

30 One type of humanized antibody is a chimera, in which the variable region genes of a non-human antibody are cloned into a human expression vector containing the appropriate human light chain and heavy chain constant region genes (72). The resulting chimeric monoclonal antibody should have the antigen-binding capacity (from the variable region of the 35 non-human source) and should be significantly less

immunogenic than the unaltered non-human monoclonal antibody.

Jones and his colleagues further humanized chimeric antibodies through a technique known as complementarity determining region (CDR) grafting (72). In this process, the antigen binding sites, which are formed by three CDRs of the heavy chain and three CDRs of the light chain, are excised from cells secreting non-human monoclonal antibodies and grafted into the DNA coding for the framework of the human antibody (72). Since only the antigen-binding site CDRs of the non-human antibody are transplanted, the resulting humanized antibody is less immunogenic than a chimeric antibody in which the entire variable domain is transplanted.

This process has been further improved by reshaping, hyperchimerization, and veneering (72). In the reshaping process on the basis of homology, the non-human variable region is compared with the consensus sequence of the protein sequence subgroup to which it belongs (72). Similarly, the selected human constant region accepting framework is compared with its family consensus sequence (72). The sequence analyses identify residues which may have undergone mutation during the affinity maturation procedure and may therefore be idiosyncratic (72). Inclusion of the more common human residues minimizes immunogenicity problems by replacing human acceptor idiosyncratic residues.

Hyperchimerization is an alternative method of identifying residues outside of the CDR regions that are likely to be involved in the reconstitution of binding activity (72). In this method, the human sequences are compared with non-human variable region sequences and the one with highest homology is selected as the acceptor framework (72). As in the reshaping procedure, the idiosyncratic residues are replaced by the more commonly occurring human residues (72). The non-

CDR residues that may be interacting with the CDR sequences are identified (72). Finally, one of these residues is selected to be included in the variable region framework (72).

5

Veneering is the process of replacing the displayed surfaces of proteins, or residues, which differ from those commonly found in human antibodies (72). Appropriate replacement of the outer residues may have little or no impact on the inner domains or interdomain framework (72). In the process of veneering, the most homologous human variable regions are selected and compared by each residue to the corresponding non-human variable regions (72). Then, the non-human framework residues, which differ from the human homologue, are replaced by the residues present in the human homologue (72).

The subject antibodies against glatiramer acetate can be administered by any method known to those of skill in the art. Such methods include, but are not limited to, intravenous, subcutaneous, intramuscular and intraperitoneal injection, and oral, nasal and rectal administration of the active substance and a pharmaceutically acceptable carrier.

25 In addition to in vivo methods of promoting remyelination, ex vivo methods of stimulating remyelination in CNS axons are encompassed by the present invention. For example, antibodies against glatiramer acetate may be used in vitro to stimulate the proliferation and/or differentiation of glial 30 cells, such as oligodendrocytes. These exogenous glial cells can then be introduced into the CNS using known techniques. Remyelination of CNS axons would be increased by raising the number of endogenous glial cells, as these cells play a critical role in the production of myelin.

35

In vitro methods of producing glial cells, or stimulating the proliferation of glial cells from mixed culture are also encompassed by the subject invention. For example, cells obtained from rat optic nerve, or rat brain, containing glial cells, are cultured as a mixed culture under conditions sufficient to promote growth of the cells. An effective amount of antibodies against glatiramer acetate is then added to the mixed culture and maintained under conditions sufficient for growth and proliferation of cells. The 5 antibodies against glatiramer acetate stimulate the proliferation of glial cells in the mixed culture. Thus, the proliferation of glial cells cultured in the presence of antibodies against glatiramer acetate is increased, relative to the proliferation of glial cells grown in the absence of 10 the antibodies. 15

The subject invention concerns treatment of a demyelinating central nervous system disease by a treatment regime that promotes production of a level of antibodies against 20 glatiramer acetate or polyclonal B cell expansion that results in CNS remyelination.

In one embodiment, the treatment regime entails the administration of glatiramer acetate. Regime variables could 25 include, but not be limited to, dose, frequency of administration, sites of administration, and adjuvant co-administration. The appropriate regimen of treatment with glatiramer acetate should be determined empirically from patient studies.

30

Glatiramer acetate and antibodies against glatiramer acetate can be formulated into pharmaceutical compositions containing a pharmaceutically acceptable carrier. As used herein, pharmaceutically acceptable carrier includes any and all 35 solvents, dispersion media, adjuvants, coatings,

antibacterial and antifungal agents, isotonic and absorption delaying agents, sweeteners and the like. The pharmaceutically acceptable carriers may be prepared from a wide range of materials including, but not limited to, 5 flavoring agents, sweetening agents and miscellaneous materials such as buffers and absorbents that may be needed in order to prepare a particular therapeutic composition. The use of such media and agents with pharmaceutically active substances is well known in the art. Except insofar as any 10 conventional media or agent is incompatible with the active ingredient, its use in the therapeutic compositions is contemplated.

Glatiramer acetate and antibodies against glatiramer acetate 15 can be formulated into any form known in the art using procedures available to one of skill in the art. In one embodiment, glatiramer acetate or an antibody against glatiramer acetate is introduced into the body by way of ingestion or inhalation. For example, they may be 20 administered by way of the mouth through feeding, through a stomach tube, by inhalation into the bronchial passages or by nasal inhalation. The composition contemplated by the subject invention may be administered either as a simple oral solution, as an emulsion or suspension formulation, as a 25 solid oral dosage form (capsule or tablet), or even as a soft gelatin capsule. The present invention contemplates immediate release dosage forms and modified release dosage forms (including particulates, coated granules and pellets, emulsions, microemulsions and encapsulation in microspheres 30 and nanospheres).

In one embodiment, the composition is formulated into a capsule or tablet using techniques available to one of skill in the art.

In another embodiment, glatiramer acetate or an antibody against glatiramer acetate is administered in another convenient form, such as an injectable solution or suspension, a spray solution or suspension, a rectal 5 suppository, a lotion, a gum, a lozenge, a food or snack item. Food, snack, gum or lozenge items can include any ingestible ingredient, including sweeteners, flavorings, oils, starches, proteins, fruits or fruit extracts, vegetables or vegetable extracts, grains, animal fats or 10 proteins. Thus, the present compositions can be formulated into cereals, snack items such as chips, bars, gum drops, chewable candies or slowly dissolving lozenges.

For both glatiramer acetate and antibodies against glatiramer acetate, one of skill in the art can readily substitute structurally-related amino acids without deviating from the spirit of the invention. The present invention includes polypeptides and peptides which contain amino acids that are structurally related to tyrosine, glutamic acid, alanine or lysine and possess the ability to stimulate the production of polyclonal antibodies against them. Such substitutions retain substantially equivalent biological activity in their ability to suppress or alleviate the symptoms of the CNS disease. These substitutions are structurally-related amino 20 acid substitutions, including those amino acids which have about the same charge, hydrophobicity and size as tyrosine, glutamic acid, alanine or lysine. For example lysine is structurally-related to arginine and histidine; glutamic acid is structurally-related to aspartic acid; tyrosine is structurally-related to serine, threonine, phenylalanine and tryptophan; and alanine is structurally-related to valine, leucine and isoleucine. These and other conservative 25 substitutions, such as structurally-related synthetic amino acids, are contemplated by the present invention.

Moreover, glatiramer acetate can be composed of l- or d- amino acids. As is known by one of skill in the art, l-amino acids occur in most natural proteins. However, d- amino acids are commercially available and can be substituted for some or all 5 of the amino acids used to make glatiramer acetate. The present invention contemplates glatiramer acetate consisting essentially of l-amino acids, as well as glatiramer acetate consisting essentially of d-amino acids.

10 Experimental Details

EXPERIMENTAL METHODS

Mice and Viral Infection

15 All mice used in the experiments were SJL/J mice (Jackson Laboratories, Bar Harbor, ME). Six- to eight-week-old mice were intracerebrally injected with 2.0×10^6 pfu of Daniel's strain of Theiler's murine encephalomyelitis virus.

20 Virus

The Daniel's strain of Theiler's murine encephalomyelitis virus (TMEV) was used in all experiments. The original virus stock was obtained from J. Lehrich and B. Arnason after eight passages in cultured baby hamster kidney (BHK) cells (33). 25 The virus was passaged an additional six times at a multiplicity of infection of 0.1 plaque-forming units per cell. Cell-associated virus was released by freeze-thawing and sonication. The lysate was clarified by centrifugation and stored in aliquots at -70°C.

30

Glatiramer Acetate Injections

All injections of glatiramer acetate, alone or in IFA, were given subcutaneously in the flank or neck. Glatiramer acetate was administered in an emulsion consisting of equal 35

volumes of glatiramer acetate in PBS and IFA (DIFCO, Fisher, Pittsburgh, PA).

Spinal Cord Lesions: Demyelination and Remyelination

5 Quantitation

Areas of spinal cord demyelination (lesion load) and remyelination (73) were determined from multiple cross-sections of plastic-embedded spinal cords, using a camera lucida system and a computerized drawing tablet (39).

10 Briefly, mice were sacrificed by overdose with sodium pentobarbital and perfused by intracardiac puncture with Trumps fixative, containing 4% paraformaldehyde and 1% glutaraldehyde. Spinal cords were removed, post-fixed in osmium tetroxide, and sectioned into 1 mm blocks. Every 15 third block (10-12 blocks per spinal cord; cervical to lumbar range) was embedded in Araldite plastic. The embedded tissues were cross-sectioned at 1 μ m thickness, and the slides stained with 4% paraphenylenediamine to highlight the myelin sheaths.

20 Using a camera lucida attached to a Zeiss photomicroscope and a ZIDAS interactive digital analysis system, three parameters were measured from each slide: total white matter area, demyelinated lesion area, and remyelination area. Outline of 25 these regions were traced and the areas calculated by the computerized digital analysis system. Demyelination was expressed as the total lesion area as a percentage of total white matter area. Remyelination was expressed as the total remyelination area as a percentage of the total demyelinated 30 lesion area. The criterion for remyelination by oligodendrocytes was abnormally thin myelin sheaths. All remyelination data refers to oligodendrocyte-mediated remyelination. Occasionally, Schwann-cell mediated remyelination was observed, characterized by abnormally thick 35 myelin sheaths and nuclei juxtaposed to the myelin sheath.

Brain Pathology Scoring

Mice were sacrificed and perfused by intracardiac puncture with Trumps fixative, containing 4% paraformaldehyde and 1% glutaraldehyde. Brains were removed and post-fixed in Trumps. Each brain was sectioned coronally into three pieces by cuts through the infundibulum and optic chiasm. The pieces were then dehydrate and embedded in paraffin. Sections from each block were mounted on slides and stained with hematoxylin and eosin to identify pathology in the following brain regions: cortex, corpus callosum, hippocampus, brainstem, striatum, and cerebellum. Pathologic scores were assigned without knowledge of the experimental treatment. Each area of the brain was graded as follows:

- 0 = no inflammation
- 15 1 = minimal inflammation, confined to perivasculature
- 2 = moderate inflammation, including parenchyma infiltration, but no tissue damage
- 20 3 = intense parenchyma inflammation with minor but definite tissue damage (loss of tissue architecture, cell death, neurophagia, neuronal vacuolation)
- 4 = extensive inflammation and tissue damage.

ELISA

An indirect ELISA was performed in which serum or a purified antibody was applied to plates that were pre-coated with glatiramer acetate or other protein antigens. The antigen of interest was dissolved in 0.1 M carbonate buffer, pH 9.5, and applied at 1 μ g per well in 96-well polystrene plates. Incubation was overnight at 4°C. Plates were then rinsed with PBST (phosphate buffered saline containing 0.05% Tween 20 detergent) and incubated for 1 hr in PBSM (phosphate buffered saline containing 5% defatted milk powder). Plates were rinsed with PBST, then incubated for 4 hr at room temperature with 50X PBS. Plates were rinsed with PBST, then incubated 35 with biotinylated secondary antibodies that were raised in

goat. The secondary antibodies, diluted in PBSM 1:50, were specific to mouse IgG or mouse IgM. Incubation was for 2 hr at room temperature. Plates were rinsed in PBST, then PBS. Streptavidin-alkaline phosphatase conjugate diluted in PBS 5 was applied to the plates for 2 hr room temperature. Plates were rinsed with PBST, with a final rinse in water. The colored reaction product was produced by incubation with p-nitrophenyl phosphatase in 0.1 M carbonate buffer plus 1 mM magnesium chloride. The reaction was stopped with 0.5 N 10 sodium hydroxide. Antibody isotyping was performed using a detection kit and purified isotype standards from Zymed Laboratories (San Francisco, CA). Extrapolation of the concentration of each isotype within the antibody samples was based on the sample dilution that produced the absorbence 15 signal falling within the most linear portion of each standard isotype curve. Optical absorbency was measured at a wavelength of 405 nm.

Delayed Type Hypersensitivity (DTH)

20 Mice were injected intradermally in the ear pinna with 10 μ l (3.5 μ g) of UV-inactivated TMEV or glatiramer acetate in sterile PBS using a 27-gauge needle. Ear thickness was measured prior to injection, and 24 and 48 hr after injection.

25

Immunohistochemistry on Cultured Cells

Glial cultures (mixed or oligodendrocyte-enriched) were derived from cerebral hemispheres from 4-7-day-old Sprague-Dawley rat pups (Harlan Sprague Dawley. Indianapolis, IA), 30 maintained on poly-lysine-coated glass coverslips in DMEM medium containing 10% fetal bovine serum, and immunostained between Days 4-28 in vitro. CNS glial cultures were also derived from adult human brain biopsies (obtained from surgical correction of epilepsy). Mouse peritoneal 35 macrophages were derived by lavage, 5-8 days following

intraperitoneal injection of sterile, 3% thioglycollate solution, and maintained in RPMI medium containing 5% fetal bovine serum for 1-3 weeks.

- 5 Spinal cord sections were obtained by cryostat sectioning of frozen spinal cords (10 μ m thickness). Sections were lightly fixed in ice-cold 95% ethanol for 5 min and incubated in 10% goat serum to reduce nonspecific staining.
- 10 Application of primary antibodies in PBS buffer was performed with ice-cold solutions with culture plate on ice with the intention of staining the cell surface. Primary antibodies were applied for 30-45 min. After rinsing in PBS for 10 min, fluorophore-conjugated secondary antibodies diluted in ice-cold PBS were applied for 30 min. Cells were then rinsed with PBS for 10-15 min. Fixation with 4% paraformaldehyde occurred either once, following the final PBS rinse, or twice, just prior to secondary antibody application and following the final PBS rinse. Cells were viewed with
- 15
- 20 Olympus fluorescent microscopes.

The primary antibodies included antibodies against glatiramer acetate (4-40 μ g/ml), Normal antibodies (20 μ g/ml), anti-glial fibrillary acidic protein (GFAP, an astrocyte marker) (Dako, Carpinteria, CA), O1 (mature oligodendrocyte marker), O4 (oligodendrocyte marker), A2B5 (immature oligodendrocyte marker), 94.03 (oligodendrocyte marker), isolectin B₄, CD11b (complement receptor 3) (activated microglia and macrophage markers), rat anti-F4/80 (Serotec, Raleigh, NC), biotinylated isolectin B₄ from Bandeiraea simplicifolia (Sigma), biotinylated mouse anti-MHC Class II (clone 10.2.16), biotinylated rat anti-Fc γ III/II receptor (CD16/CD32, BD PharMingen, San Diego, CA), rat anti-myelin basic protein (82-87; Calbiochem, San Diego, CA). The secondary antibodies were anti-species IgG or IgM, raised in

goat, and fluorophore-conjugated (Jackson Immunoresearch; Vector) for direct detection or biotinylated for detection by the peroxidase method using an ABC Elite kit (Vector).

5 Glatiramer acetate antibodies and normal mouse antibodies were usually applied as biotinylated derivatives. Biotinylation was performed by 30 min incubation of purified antibodies with EZ-Link NHS-LC biotin (Pierce), followed by extensive dialysis against PBS (10,000 molecular weight 10 cutoff). Biotinylation and preservation of glatiramer acetate binding activity was confirmed by Western blot and ELISA. Purified unbiotinylated glatiramer acetate antibodies used in conjunction with a Mouse-on-Mouse Staining Kit (Vector) showed the same staining pattern as biotinylated 15 glatiramer acetate antibodies.

Purification of Antibodies Against Glatiramer Acetate in SJL/J Mice (Figure 29)

20 The steps involved in purification of antibodies against glatiramer acetate were:

1. Immunization of SJL/J mice. Mice were immunized by eight, subcutaneous injections of glatiramer acetate in IFA (0.1 mg/injection between Days 0-50).
2. Serum isolation. Mice were bled 9 times between Days 25 14-56 from the time of first immunization. After each bleed, blood was stored overnight at 4°C, then centrifuged to isolate serum. Serum was stored at -20°C until all bleeds were completed.
3. Affinity column isolation of antibodies against 30 glatiramer acetate. Glatiramer acetate was coupled to NHS-activated Hi-Trap affinity columns (Amersham Pharmacia) by the manufacturer's recommended procedures. Serum was thawed, pooled, centrifuged to remove serum lipid, diluted with phosphate buffer, and run over the 35 glatiramer acetate affinity column twice. After rinsing

the column, antibodies against glatiramer acetate were eluted with glycine-HCl (pH 3.0) into Tris buffer (pH 8.0). The OD₂₆₀ of the eluted samples was determined as a measure of protein content. Most of the antibodies against glatiramer acetate were eluted in the first 2 fractions.

5. Dialysis. Eluted fractions containing high levels of antibodies against glatiramer acetate were pooled and dialyzed in PBS (10,000 molecular weight cut-off).
- 10 5. Purity and protein analysis. Dialyzed antibodies against glatiramer acetate were analyzed by sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) and found to be almost exclusively comprised of product representative of antibody heavy and light chains. Bicinchoninic acid protein assay was used to determine the total protein content of the sample, using bovine serum albumin (BSA) as the standard. The determined concentration of 4.8 mg/ml was considered as the concentration of antibodies against glatiramer acetate.
- 15 6. Storage. Dialyzed antibodies against glatiramer acetate were filter-sterilized with a 0.22 μ m filter and stored at 4°C. Over 2 years of storage, no precipitate or contamination was observed and glatiramer acetate reactivity by ELISA and cellular binding was retained.
- 20
- 25

Purification of Normal Antibodies (Figure 30)

The steps involved in purification of Normal antibodies were:

1. Protein A/G column isolation of antibodies against glatiramer acetate. Normal mouse serum (SIGMA commercial preparation, St. Louis, MO) was centrifuged to remove serum lipid, diluted with phosphate buffer, and run over a Protein A/G column (Hi-trap Protein A/G column; Amersham Pharmacia). After rinsing the column, Normal antibodies were determined as a measure of
- 30
- 35

protein content.

2. Dialysis. Eluted fractions containing high levels of antibodies were pooled and dialyzed in PBS.

3. Purity and protein analysis. Dialyzed Normal antibodies were analyzed by sodium dodecyl sulphate-polyacrylamide gel electrophoresis (SDS-PAGE) and found to be almost exclusively comprised of product representative of antibody heavy and light chains. Bicinchoninic acid protein assay was used to determine the total protein content of the sample, using BSA as the standard. The determined concentration of 2.6 mg/ml was considered as the Normal antibody concentration.

10

4. Storage. Dialyzed Normal antibodies were filter-sterilized with a 0.22 μ m filter and stored at 4°C.

15 Over 2 years of storage, no precipitate or contamination was observed.

Normal antibodies were also purified by running Normal mouse serum (SIGMA commercial preparation, St. Louis, MO) over Protein A/G columns (Pierce, Rockford, IL). The antibodies were eluted by conditions similar to those for glatiramer acetate antibodies.

Antibody Injections

25 All antibodies were dissolved in PBS and administered intraperitoneally. Normal antibodies were protein G-purified antibodies isolated from commercially purchased mouse serum. These were used as a control for antibodies against glatiramer acetate.

30

EXAMPLE 1: EFFECT OF GLATIRAMER ACETATE TREATMENT ON EXTENT OF SPINAL CORD WHITE MATTER PATHOLOGY DURING EARLY DISEASE.

Procedure

Mice were injected with glatiramer acetate (0.1 mg/injection) in IFA or IFA alone on Day -15 and Day 7, relative to virus injection. Additional injections of either glatiramer acetate alone or phosphate buffered saline (PBS) were performed on Days -7, 0, 13, 21, 32, and 41. Virus was injected on Day 0. Mice were sacrificed on Day 45 post-infection (60 days of glatiramer acetate treatment), and spinal cord demyelinating pathology was measured.

10

Results

Glatiramer acetate exerted no statistically significant effect on the extent of demyelination by 45 days post-infection (Figure 1). However, a trend towards reduced demyelination was present (T-test, $P=0.08$).

EXAMPLE 2: EFFECT OF GLATIRAMER ACETATE TREATMENT DURING LATE DISEASE

20 Experiment 2A: Effect of Glatiramer Acetate on the Extent of Spinal Cord White Matter Pathology During Late Disease

Procedure

Chronically infected mice (124-365 days post-infection) were 25 treated subcutaneously for periods of 41-76 days. One group of mice received 0.1 mg glatiramer acetate by 8 injections of 0.1 mg each. These mice were sacrificed after 65 days. A second group of mice received 0.1 mg glatiramer acetate/IFA by 4-8 injections at equal intervals, 0.1 mg/injection. The 30 members of this group were sacrificed after 65-76 days. A third group of mice received 5 mg glatiramer acetate/IFA by 1-2 injections each of 5 mg, an effective dose for inhibiting EAE (32). Sacrifice of these mice occurred after 41-65 days. The control group of mice received PBS. In all groups, 35 spinal cord demyelination was measured after sacrifice.

Results

Treatments with the high dose of glatiramer acetate caused an expansion of the demyelinating lesions ($P < 0.05$, compared to PBS treated mice, unpaired t-test) (Figure 2). Low doses of 5 glatiramer acetate, either alone or in combination with adjuvant, did not alter the extent of demyelination. Lesions were characterized by extensive primary demyelination and infiltration with macrophages and lymphocytes (Figure 21). On average, 10% of the white matter was demyelinated after 10 treatment with PBS, IFA, or low-dose glatiramer acetate. In contrast, the mean lesion load doubled to 20% after immunization with a high dose of glatiramer acetate, which was significant by t-test ($P=0.01$ vs IFA) and by one-way analysis of variance (ANOVA, $P=0.006$, comparing all groups).

15

Experiment 2B: Effect of Glatiramer Acetate on Extent of Remyelination in the Spinal Cord During Late Disease

Procedure

20 Chronically infected mice (124-365 days post-infection) were treated subcutaneously with glatiramer acetate as in Experiment 2A. After sacrifice, spinal cord remyelination was measured in all mice. Spinal cords with less than 4.0% demyelination were excluded from the remyelination analysis 25 (2 from PBS, 1 from IFA and 2 from 0.1 mg glatiramer acetate).

Results

Treatment with glatiramer acetate alone or with IFA did not 30 affect the extent of oligodendrocyte-mediated remyelination during late disease (Figure 3). Spontaneous remyelination in PBS-treated mice represented 8.5% of the lesion area. IFA alone enhanced remyelination ($p < 0.05$, compared to PBS, unpaired t-test). The beneficial effects of IFA raises the 35 possibility that polyclonal B cell activation with

appropriate stimulants might promote remyelination. No treatment effect was seen on the extent of Schwann cell-mediated remyelination, characterized by abnormally thick myelin sheaths, adjacent Schwann cell bodies, and more widely dispersed axonal profiles.

Experiment 2C: Effect of Glatiramer Acetate on Brain Pathology During Late Disease

10 Procedure

Chronically infected mice (250 days post-infection) were treated by subcutaneous injection with either PBS, IFA or 1.0 mg glatiramer acetate with IFA, twice weekly for a total of 8 injections. After 29 days of treatment, mice were 15 sacrificed and paraffin-embedded brain sections were scored for the extent of pathology, as explained in the Experimental Methods section.

20 Results

Figure 4 shows the effect of glatiramer acetate on brain pathology. Each symbol is the score from an individual animal. Glatiramer acetate treatment did not affect the extent of the brain pathology during late disease.

25 Experiment 2D: Effect of High Dose Glatiramer Acetate on Anti-viral Immunity

Procedure

Chronically infected mice (124-365 days post-infection) were 30 treated subcutaneously with glatiramer acetate as in Experiment 2A. Serum titers of antibodies against TMEV and anti-TMEV DTH reactions were measured.

Results

Four weeks after glatiramer acetate immunization, anti-TMEV titers were reduced (total IgG, IgG1, and IgG2a), compared to control mice, which were statistically significant in 6 of 12 dilutions among the 3 classes of antibodies (Table 1).

Table 1. Changes in Anti-viral Immunity Following Glatiramer Acetate Immunization

10

<u>Anti-TMEV IgG ^{a,d}</u>		<u>TMEV DTH ^{a,c}</u>		<u>TMEV Antigen Expression ^b</u>	
	OD	n	Δ mm	n	#of cells/mm ²
15	Control 1.08 ± 0.15 ^e		0.125		9.01 ±
	5mg GA/IFA 0.69 ± 0.05 ^g	10	± 0.024 ^e	11	1.71 ^f
		11	0.061	10	10.83 ± 0.86
			± 0.039 ^h		

All values are means ± SEM

20 n = number of mice

^a 4 wk post-immunization

^b 6-9 wk post-immunization

^c measured 48 hr post-injection of UV-TMEV

^d 6250X serum dilution

25 ^e IFA control immunization

^f PBS control immunization

^g p<0.05 vs IFA, unpaired t-test

^h p<0.05 vs PBS, Mann Whitney test

30 Anti-TMEV DTH reactions were also reduced in immunized mice. In 9 out of 10 immunized mice, the TMEV DTH was below the

mean of the IFA control group. The remaining mouse had an inexplicably high response that exceeded all control mice. As expected from reduced antiviral immunity, quantitation of virus antigen-positive cells revealed that virus expression 5 tended to be higher 6-9 wks after high-dose immunization. In the glatiramer acetate group, virus expression in 7 out of 8 mice exceeded the mean expression in the control group. As in the DTH experiments, 1 of 12 mice in the PBS control group had uncharacteristically high virus expression (6 standard 10 deviations from the mean of the remaining 11 mice). These results show that glatiramer acetate at a high dose mildly reduced cellular and humoral immunity to TMEV. This reduced antiviral immunity, resulting in increased viral pathogenesis, may account for the increase in lesion load. 15 The reduction may have resulted from the effects of glatiramer acetate-specific T cells which were found in both infected and uninfected mice.

Experiment 2E: Effect of Antibodies Against Glatiramer Acetate on Demyelination During Late Disease

Procedure

Chronically infected mice (160-477 days post-infection) were treated by intraperitoneal injection with antibodies for 25 periods of 36-76 days. One group of mice received Normal antibodies, administered in 10 injections of 0.05 mg each for a total of 0.5 mg over 42 days. A second group of mice received 0.5 mg antibodies against glatiramer acetate, administered in 10 injections of 0.05 mg each, for a total of 30 0.5 mg over 41 - 42 days. The third group of mice received 1.5 mg antibodies against glatiramer acetate, administered in 5 injections of 0.3 mg each for a total of 1.5 mg over 36 days. PBS was administered to the control group of mice. Spinal cord white matter demyelinating pathology was measured 35 post-sacrifice for all mice.

Results

Neither antibodies against glatiramer acetate nor Normal antibodies, when passively transferred, affected the extent
5 of demyelinating pathology during late disease (Figure 5).

Experiment 2F: Effect of Antibodies Against Glatiramer Acetate on Demyelination During Late Disease

10 (i)

Procedure

Chronically diseased, virus-infected mice were treated for 5-
6 wks with glatiramer acetate antibodies or normal
antibodies. Each mouse received 5 weekly injections for
15 total antibody doses of 0.5 or 1.5 mg (0.1 - 0.3
mg/injection). After sacrifice, spinal cord demyelination
was measured.

20 Results

None of the antibody treatments affected lesion load, as
compared to PBS-treated mice (Figure 7a), indicating that
glatiramer acetate antibodies were not pathogenic at serum
levels of approximately 200 μ g/ml or less.

25

(ii)

Procedure

Chronically infected mice (160-477 days post infection) were
treated by antibodies or PBS, following the procedure of
30 Experiment 2E(i). After sacrifice, spinal cord remyelination
was measured. Spinal cords with less than 4.0% demyelination
were excluded from the remyelination analysis (2 from PBS, 2
from 0.5 mg antibodies against glatiramer acetate, and 1 from
1.5 mg antibodies against glatiramer acetate).

35

Results

Glatiramer acetate antibody treatment at both doses increased the extent of remyelination ($P < 0.05$ for both compared to PBS, unpaired t-tests) (Figure 6). Normal antibodies did not 5 affect remyelination, suggesting that the beneficial effect of antibodies against glatiramer acetate was through specific antigen (epitope) interactions rather than through nonspecific interactions of antibody heavy chains with F_c receptors.

10

Experiment 2G: Effect of Antibodies Against Glatiramer Acetate on Remyelination During Late Disease

Procedure

15 Chronically diseased, virus-infected mice were treated for 5-6 wks with glatiramer acetate antibodies or normal antibodies. Each mouse received 5 weekly injections for total antibody doses of 0.5 or 1.5 mg (0.1 - 0.3 mg/injection). Spinal cord remyelination was measured after 20 sacrifice.

Results

Glatiramer acetate antibodies had a positive therapeutic effect on oligodendrocyte-mediated remyelination (Figure 7b). 25 At a total dose of 1.5 mg, mean remyelination was increased 2.4-fold by glatiramer acetate antibodies, which was significant by t-test ($P=0.02$ vs PBS) and one-way ANOVA ($P=0.04$ comparing PBS, SJL Ig, pooled Ig, and 1.5 mg glatiramer acetate Ig). A positive treatment effect was also 30 indicated by a strong statistical difference between the variances of the 1.5 mg glatiramer acetate antibody group and the other treatment groups (Bartlett's test, $P=0.003$). Based on axon densities within remyelinated regions, the 2.4-fold increase in remyelination was equivalent to the repair of 35 approximately 7,000-21,000 internodes in the sections that we

analyzed. Remyelination also tended to increase with 0.5 mg glatiramer acetate antibody treatment but without statistical significance. Qualitatively, glatiramer acetate antibody-promoted remyelination was of the highest quality seen in 5 these experiments, with some lesions being nearly completely repaired (Figure 28). Neither normal SJL immunoglobulin nor pooled mouse immunoglobulin significantly promoted remyelination. Thus, in the absence of complete glatiramer acetate immunization and in the absence of oligodendrocyte 10 binding, antibodies to glatiramer acetate stimulated myelin repair.

EXAMPLE 3: SERUM TITRES OF GLATIRAMER ACETATE IgG

15 Experiment 3A: Serum Titres of Glatiramer Acetate IgG in Non-infected, Immunized Mice

Procedure

Non-infected mice were immunized with glatiramer acetate/IFA 20 on Days 0, 4, 8, 15, and 26 (0.1 mg glatiramer acetate/injection) or on Days 0, 3, 7, 10, and 21 (1.0 mg/injection). On Days 0, 7, 14, 19, 28, 33 and 51, blood was collected from 26-35 mice, serum was isolated by glatiramer acetate affinity chromatography (GA Ig) and 25 pooled. Antibodies from normal SJL serum (SJL Ig) and commercial, pooled mouse serum (Pooled Ig) were also purified by Protein A/G chromatography. ELISA was performed using glatiramer acetate-coated plates and biotinylated anti-mouse IgG as the secondary antibody.

30

Results

Shown in Figure 8 are the 1:1000 sera dilutions. Serum titres of glatiramer acetate IgG were first detectable 14 days post-immunization and increased over time.

35 Approximately 3 weeks were required to achieve high antibody

titres. Varying the glatiramer acetate doses from 0.1 - 1.0 mg and altering the timing of injections did not significantly influence glatiramer acetate IgG titres, although the immunization and sampling regimens also 5 differed. These data helped to determine how long infected mice should be treated with glatiramer acetate in order to evaluate whether antibodies against glatiramer acetate generated by immunization of infected mice can promote remyelination.

10

Experiment 3B: Serum Titres of Glatiramer Acetate IgG in Chronically Diseased Mice After Immunization

Procedure

15 Mice infected for 124 days were immunized with glatiramer acetate or glatiramer acetate/IFA at 0.1 mg/injection for a total for 0.8 mg from Days 0-50. Mice were sacrificed on Day 65 post-immunization. Then, serum was isolated and ELISA was performed on glatiramer acetate-coated plates. The 20 secondary antibody was biotinylated anti-mouse IgG.

Results

Serum titres of glatiramer acetate IgG were first detectable 25 14 days following immunization and continued to rise thereafter. There was significant variability in the production of antibodies against glatiramer acetate among individual mice immunized with glatiramer acetate alone (Figure 9). In contrast, antibody levels were more similar 30 among mice treated with glatiramer acetate and IFA. Immunization with glatiramer acetate and IFA generated much higher antibody levels than immunization with glatiramer acetate alone. No glatiramer acetate IgG was detected in serum from mice treated with either PBS or IFA alone.

35

Experiment 3C: Serum Titres of Glatiramer Acetate IgG in Chronically Diseased Mice

Procedure

5 Chronically diseased mice, infected for 196-286 days, were immunized with glatiramer acetate. The first group of mice received 5 mg glatiramer acetate/IFA injections on Days 0 and 20. Serum was isolated from these mice on Day 62. The next group of mice received 5 mg glatiramer acetate/IFA injections
10 on Day 0. Serum was isolated from these mice on Day 41. The last group of mice received 0.1 mg glatiramer acetate/IFA injections on Days 0, 25, 37, and 64. Serum was isolated on Day 76. For all groups, ELISA was performed using glatiramer acetate-coated plates. The secondary antibody was
15 biotinylated anti-mouse IgG.

Results

Again, serum titres of glatiramer acetate IgG were first detectable 14 days following immunization and increased
20 thereafter. Antibody titers reached a high of 1.2 mg/ml in mice immunized with 5 mg glatiramer acetate (Figure 12a). The production of antibodies against glatiramer acetate was highly dose-dependent (Figure 10). Very high glatiramer acetate IgG titres were reached after immunization and a
25 single boost using 5 mg glatiramer acetate injections. The titres were much higher than following 4 injections of 0.1 mg glatiramer acetate. However, relatively high titres were reached even using 0.1 mg glatiramer acetate over time. Individual symbols are the means (\pm SEM) for 4-5 mice, except
30 for the untreated group (2 mice).

Following Treatment with Passively Transferred Antibodies Against Glatiramer Acetate vs. Glatiramer Acetate Immunization

5 Procedure

One group of chronically diseased mice (196-348 days of infection) received 5 mg glatiramer acetate in IFA in a single subcutaneous injection. The other group of chronically diseased mice received 10 x 50 µg intraperitoneal 10 injections of antibodies against glatiramer acetate, twice weekly. The final passive transfer was on Day 35. After 41 days of treatment, both groups of mice were sacrificed and serum titres of Copolymer IgG was measured by ELISA. The secondary antibody was biotinylated anti-mouse IgG.

15

Results

Glatiramer acetate IgG was detected in serum 6 days following the final passive transfer of antibodies against glatiramer acetate, but levels were much lower than in mice that received a single immunization with high dose glatiramer acetate (Figure 11). Since IgG is cleared with a half-life of approximately 3 weeks, the low serum titre of antibodies against glatiramer acetate 6 days after the final passive transfer suggests that low levels of antibodies against 20 glatiramer acetate are sufficient for promotion of remyelination.

EXAMPLE 4: REACTIVITY OF PURIFIED ANTIBODIES AGAINST GLATIRAMER ACETATE AND PURIFIED NORMAL ANTIBODIES

30

Experiment 4A: Glatiramer Acetate Reactivity of Purified Antibodies Against Glatiramer Acetate and Purified Normal Antibodies: IgG and IgM

35 Procedure

The procedures for isolating antibodies against glatiramer acetate and Normal IgG were as described in Experimental Methods. Purified antibodies against glatiramer acetate or Normal antibodies were assayed by ELISA using glatiramer acetate-coated plates. The secondary detection antibodies were either biotinylated anti-mouse IgG or biotinylated anti-mouse IgM.

Results

10 Antibodies against glatiramer acetate had high reactivity to glatiramer acetate (Figure 13). Both glatiramer acetate IgG and glatiramer acetate IgM were detected. Normal antibodies had little or no IgG- or IgM-reactivity to glatiramer acetate.

15 Experiment 4B: Protein Polyreactivity of Purified Antibodies Against Glatiramer Acetate and Purified Normal Antibodies

Procedure

20 Proteins were adsorbed to ELISA plates as described in Experimental Methods. Plates were then reacted with antibodies against glatiramer acetate, Normal antibodies, or SCH 94.03 monoclonal IgM antibodies. Secondary antibodies were anti-mouse IgG or anti-mouse IgM.

25

Results

Very low polyreactivity of glatiramer acetate IgG/IgM and Normal IgG was observed (Figure 14). The only significant cross-reactivity was to rabbit myosin heavy chain. This 30 contrasts with multiple cross-reactivities seen for SCH 94.03 IgM, as reported in U.S. Patent 5,591,629 (1). This supports the hypothesis that the mechanism by which antibodies against glatiramer acetate promote remyelination is different from the mechanism used by polyreactive IgM antibodies such as SCH 35 94.03.

EXAMPLE 5: ISOTYPE ANALYSIS OF ANTIBODIES AGAINST GLATIRAMER ACETATE AND NORMAL ANTIBODIES

5 Procedure

Purified antibodies against glatiramer acetate and Normal antibodies were isotyped by ELISA using an antibody isotyping kit (Pierce).

10 Results

All tested isotypes were found in both glatiramer acetate antibodies and Normal antibodies (Figure 15). Glatiramer acetate IgG reactivity in the polyclonal glatiramer acetate Ig preparation was strong, being detectable by enzyme-linked immunoabsorbent assay (ELISA) at a concentration as low as 8 ng/ml, whereas IgM reactivity was barely detectable. The glatiramer acetate antibody preparation had a higher relative level of IgG1 than other isotypes, compared to the IgG1 level in Normal antibodies. No glatiramer acetate-reactive IgG or IgM was found in either of the control Ig preparations. By extrapolation from standard curves of purified immunoglobulin isotypes, it was determined that IgG1 comprised 70% of the purified glatiramer acetate Ig (Figure 12b). IgG2b was the next abundant, representing 18% of the pool. In contrast, IgG2b was most abundant in purified SJL Ig (45%) and IgA was most abundant in the pooled mouse Ig (32%). These results indicated that glatiramer acetate antibodies were produced in abundance in immunized mice, either infected or uninfected, and consisted primarily of isotypes known to poorly activate complement in mice. IgG1 (non-complement fixing in mice) appears to be the most abundant in the antibodies against glatiramer acetate. The apparent abundance of IgG1 implies that complement activation would not be a significant consequence of glatiramer acetate antibody treatment.

EXAMPLE 6: EFFECT OF ANTIBODIES AGAINST GLATIRAMER ACETATE ON
IN VITRO PROLIFERATION OF LYMPH NODE-DERIVED LYMPHOCYTES FROM
MBP₈₄₋₁₀₂ AND PLP₁₇₉₋₁₉₁ IMMUNIZED MICE

5 Experiment 6A: Antibodies Against Glatiramer Acetate
Stimulate in Vitro Proliferation of Lymph Node-derived
Lymphocytes from Mice

Procedure

10 Two mice were subcutaneously injected in the flanks with MBP₈₄₋₁₀₂ in complete Fruend's adjuvant (CFA). MBP₈₄₋₁₀₂ was obtained by following the procedure of Hawes et al (24). A method similar to that of Tuohy et al. was employed to produce PLP₁₇₉₋₁₉₁ (70). PLP₁₇₉₋₁₉₁ in CFA was subcutaneously
15 injected into the flanks of 1 mouse. After 10 days, inguinal and per-aortic lymph nodes were removed, dissociated, and grown for 53 hours in culture in medium alone or in the presence of MBP₈₄₋₁₀₂, PLP₁₇₉₋₁₉₁, Normal antibodies, antibodies against glatiramer acetate or glatiramer acetate. In the
20 last 13 hours of culture, 1 μ Ci of [³H]-thymidine was added and its incorporation measured by scintillation counting of harvested cells.

Results

25 Antibodies against glatiramer acetate (25 μ g/ml) stimulated lymphocyte proliferation to a comparable level as 50 μ g/ml of specific peptide (stimulation indices = 12-17) (Figure 16). Normal antibody-induced proliferation occurred, but to a much lower level than that induced by antibodies against
30 glatiramer acetate or a specific peptide, suggesting that antigen-specificity of antibodies against glatiramer acetate contributed to its effect. Glatiramer acetate did not induce significant proliferation. The stimulation of proliferation is consistent with an effect of antibodies against glatiramer acetate on antigen presentation, but other explanations, such

as direct binding to lymphocytes, are also possible.

Experiment 6B: Antibodies Against Glatiramer Acetate Stimulate in Vitro Proliferation of Lymph Node-derived Lymphocytes from MBP₈₄₋₁₀₂-immunized Mice

Procedure

Mice were subcutaneously injected in the flanks with MBP₈₄₋₁₀₂ (2 mice) peptide in CFA. After 10 days, inguinal and peri-aortic lymph nodes were removed, dissociated, and grown for 74 hours in cultures in medium alone or with additional additives as shown in Figure 17. In the last 12 hours of culture, 1 μ Ci of [³H]-thymidine was added and its incorporation measured by scintillation counting of harvested cells.

Results

Antibodies against glatiramer acetate (25 μ g/ml) stimulated lymphocyte proliferation (stimulation indices = 6 and 13) (Figure 17). Normal antibodies did not induce proliferation at a comparable antibody concentration. As in Experiment 6A, the stimulation of proliferation could be attributed to the effect of antibodies against glatiramer acetate on antigen presentation, but there are other viable theories, such as direct binding to lymphocytes.

Experiment 6C: Antibodies Against Glatiramer Acetate Do Not Stimulate in Vitro Proliferation of Lymph Node-derived Lymphocytes from Non-immunized Mice

30

Procedure

Inguinal and peri-aortic lymph nodes from non-immunized SJL/J mice were removed, dissociated, and grown for 74 hours in culture in medium alone or in the presence of MBP₈₄₋₁₀₂, PLP₁₇₉₋₁₉₁, Normal antibodies, antibodies against glatiramer

acetate or glatiramer acetate. In the last 12 hours of culture, 1 μ Ci of [3 H]-thymidine was added and its incorporation measured by scintillation counting of harvested cells.

5

Results

Figure 18 shows that no treatment, including antibodies against glatiramer acetate (25 μ g/ml), stimulated lymphocyte proliferation to a significant degree over the baseline level (medium alone). This suggests that antibodies against glatiramer acetate stimulate lymphocyte proliferation (Figures 16 - 17) only during an active immune response to myelin peptides or other antigens.

15 EXAMPLE 7: GLATIRAMER ACETATE ANTIBODY BINDING TO CULTURED CELLS

Experiment 7A: Glatiramer Acetate Antibody Binding to Cultured Cells from the Central Nervous System

20

Procedure

The methodology is described above in the Experimental Methods. Briefly, all staining was performed with ice-cold solutions, with the culture plate on ice, and prior to fixation in order to bind the cell surface. The primary antibodies for these experiments included antibodies against glatiramer acetate, 4-40 μ g/ml, Normal antibodies, 20 μ g/ml, anti-GFAP (astrocyte markers), O1, O4, A2B5, 94.03 (oligodendrocyte markers), isolectin B₄, CD11b (complement receptor 3) (activated microglia and macrophage markers). The secondary antibodies were directed against IgG or IgM of the appropriate species.

Results

CNS glial cultures derived from neonatal rat brains

Glatiramer acetate IgG and IgM (secondary antibodies were isotype-specific) stained a small population of cells that were distinct from oligodendrocytes. Mature oligodendrocytes were readily identified by elaborate process extension and by staining with oligodendrocyte markers. In contrast, glatiramer acetate antibody-positive cells did not have elaborate process extension. Rather, they had the phenotype of activated microglia (60) as they were round, located only on the top surface of the culture, sometimes in clusters and always positive with the microglia/macrophage marker, *Bandeiraea simplicifolia* isolectin B₄ (Figure 23a). They were easily distinguished from early lineage oligodendrocytes (A2B5-positive; Figure 23b), differentiated oligodendrocytes (O1-, O4-, or MBP-positive; Figure 23c), astrocytes (glial fibrillary acidic protein (GFAP)-positive) and activated microglia which adhered to the coated-glass substrate. No co-labeling of cells was observed with antibodies against glatiramer acetate or any of the oligodendrocyte markers, whereas cells were co-labeled with glatiramer acetate IgG and the activated microglia markers, isolectin B₄ or Mac-1.

Stronger staining by antibodies against glatiramer acetate was observed after using secondary antibodies directed against mouse IgG than against mouse IgM. Immunostaining with antibodies against glatiramer acetate exceeded the staining by Normal IgG and by secondary antibodies alone, suggesting that antibodies against glatiramer acetate recognized specific cell-surface epitopes rather than being bound simply by non-specific F_c receptors. Very little immunostaining of GFAP (an intracellular antigen) was observed, suggesting that the staining seen with other antibodies (i.e., antibodies against glatiramer acetate) was

against cell surface antigens.

Human Mixed Glial Cultures (Figure 20)

In CNS glial cell cultures, a population of cells stained 5 positive for glatiramer acetate IgG. These cells were similar in phenotype to those in rat glial cultures and distinct from the O4-positive oligodendrocytes, which had elaborate process extension. These data are consistent with glatiramer acetate IgG staining of activated microglia, 10 rather than oligodendrocytes.

Experiment 7B: Binding to Cultured Mouse Peritoneal Macrophages by Antibodies Against Glatiramer Acetate

15 Procedure

The methodology is described above in the Experimental Methods. Briefly, peritoneal macrophages (similar in phenotype and function as activated microglia) were derived from the peritoneum of SJL mice, 5 days after intraperitoneal 20 stimulation with 3% sterile thioglycollate broth. Cells were then cultured for 1-3 weeks prior to staining. Antibodies were diluted in ice-cold PBS and the solutions applied to culture plates on ice in order to detect cell surface staining. The primary antibody incubations consisted of 25 combinations of the following: antibodies against glatiramer acetate, 40 μ g/ml, Normal antibodies, 20 μ g/ml, isolectin B₄, CD11b (complement receptor 3) (activated microglia and macrophage markers), glatiramer acetate, 80-200 μ g/ml.

30 Results

Figure 19 demonstrates that incubation with antibodies against glatiramer acetate under cold, unfixed conditions resulted in IgG staining of a subset of cultured mouse 35 peritoneal macrophages, which is consistent with binding to a microglial lineage cell. Surface binding in both glial and

macrophage cultures was not simply due to nonspecific interaction with Fc receptors, as no staining was observed with pooled mouse Ig. Acetone fixation/permeabilization prior to incubation with antibodies against glatiramer acetate resulted in intense staining of all cells, suggesting that antibodies against glatiramer acetate were highly reactive to intracellular antigens. Normal antibodies did not stain acetone-fixed cells. Fixation with 4% paraformaldehyde prior to incubation with antibodies against glatiramer acetate did not significantly alter the staining pattern as compared to fixation following incubation with antibodies against glatiramer acetate. Glatiramer acetate IgG-positive cells always co-stained with isolectin B₄ and Mac-1.

15

Four-hour pre-incubation of macrophages at 37°C with a variety of agents (20 µg/ml glatiramer acetate, myelin homogenate, kidney homogenate, MBP₈₄₋₁₀₂, PLP₁₇₉₋₁₉₁) did not influence subsequent staining by glatiramer acetate IgG, compared to cells preincubated only with medium. This suggests that MHC Class II presentation of processed glatiramer acetate or other antigens at the cell surface did not influence the binding of antibodies against glatiramer acetate.

25

Co-incubation of antibodies against glatiramer acetate with glatiramer acetate and macrophages greatly increased the intensity of glatiramer acetate IgG staining and the number of cells stained, suggesting that antibodies against glatiramer acetate complexed with glatiramer acetate bound much more extensively to macrophages than antibodies against glatiramer acetate alone. Co-incubation of antibodies against glatiramer acetate with myelin homogenate, SCH, or kidney homogenate did not alter glatiramer acetate staining. Punctate staining was present only when antibodies against glatiramer acetate were co-incubated with glatiramer acetate.

This staining pattern was indicative of clustered receptors, suggesting that MHC Class II molecules may have been bound by glatiramer acetate.

5 Glatiramer acetate IgG (biotinylated) bound extensively to spinal cord sections from both normal and Theiler's virus-infected mice (Figure 23d). Similar to glial cultures, reactivity to oligodendrocytes or CNS myelin was not observed. In the white matter, glatiramer acetate IgG 10 recognized a subset of presumed glia, particularly their network of thin, randomly oriented processes. Glatiramer acetate Ig-positive structures often co-labeled with GFAP, indicative of astrocyte recognition. Glatiramer acetate IgG also outlined most perivascular infiltrating cells in 15 lesioned cords, which consist of macrophages, microglia, and lymphocytes. Only a low level of background staining was found using control biotinylated pooled mouse Ig (Figure 23e).

20 Discussion

By morphological and co-immunolabeling criteria, antibodies against glatiramer acetate bound to subpopulations of activated microglia and macrophages in culture. This staining pattern contrasts with that of monoclonal antibody 25 SCH 94.03 and other remyelination-promoting antibodies, which bind preferentially to the surface of oligodendrocytes (1). Glatiramer acetate IgG staining was stronger than glatiramer acetate IgM staining. Antibodies against glatiramer acetate bound both surface and intracellular antigens. Staining of 30 antibodies against glatiramer acetate was greatly increased by co-incubation with glatiramer acetate, but not by pre-incubation of macrophages with glatiramer acetate.

35 The increased staining following co-incubation with glatiramer acetate might reflect binding of glatiramer

acetate: antibody complexes to MHC Class II molecules, which are known to bind glatiramer acetate (19, 69). In addition, the binding to microglia/macrophages in vivo might modulate cellular function, thereby triggering a more permissive environment for remyelination.

EXAMPLE 8: LESION PATHOLOGY AND REMYELINATION

Procedure

10 Chronically infected mice (6 months or longer post-infection) were treated with PBS, glatiramer acetate (0.1 mg/injection), or antibodies against glatiramer acetate (1.5 mg total).

Results

15 PBS Treatment

Mice treated with PBS showed extensive demyelination and macrophage filtration of lesions. Remyelination, characterized by abnormally thin myelin sheaths, was virtually absent (Figures 21 - 22).

20

Glatiramer Acetate Treatment

Lesions were extensively demyelinated and infiltrated with macrophages in glatiramer acetate-treated mice (Figures 25 - 26). Patches of significant remyelination were occasionally observed in some lesions, but quantitatively, the remyelination was not more extensive than following the PBS treatment.

Treatment with Antibodies Against Glatiramer Acetate

30 The lesions of mice treated with antibodies against glatiramer acetate showed extensive oligodendrocyte-mediated remyelination (Figures 27 - 28).

DISCUSSION

35 Glatiramer acetate has proved to be effective in treating multiple sclerosis (2-6, 31). Due to several inhibitory

properties, glatiramer acetate can be classified as a mixture of latered peptide ligands, an area of major interest in multiple sclerosis research (13, 28). No prior studies have proposed that glatiramer acetate works by inducing the 5 production of polyclonal antibodies against glatiramer acetate. Instead, scientists have theorized that glatiramer acetate disrupts the MHC/TCR complex formation to specific peptides (51), induces glatiramer acetate-specific suppressor cells in vivo (9) or binds directly to major 10 histocompatibility complex class II to replace MBP peptides (5, 69).

Without being limited to any specific mechanism, one hypothesis regarding the mechanism of glatiramer acetate in 15 the treatment of CNS diseases is that it actively induces a protective humoral immune response. In glatiramer acetate-treated multiple sclerosis patients, a Th2 cytokine shift in glatiramer acetate-reactive lymphocytes occurs, which is consistent with the generation of suppressor lymphocytes (17, 20 45, 50). Preliminary data indicate that multiple sclerosis patients treated with glatiramer acetate develop very high antibody titers to glatiramer acetate. There seems to be a strong positive correlation between the presence of antibodies against glatiramer acetate and therapeutic 25 efficacy. Another indication that glatiramer acetate stimulates the immune system is evident in the localized swelling and rare hypersensitivity reactions in response to glatiramer acetate. Since the most prevalent aim in multiple sclerosis therapy is to temper immune activity, it is 30 unexpected that a compound that elicits strong immune responses is therapeutic.

One theory concerning the way in which antibodies promote 35 remyelination is that by binding to oligodendrocytes or their progenitors, the antibodies may directly stimulate

proliferation or migration of oligodendrocytes, or differentiation of oligodendrocyte progenitors. It is possible to promote remyelination only after approximately 4 months of infection, a time at which most of the active 5 myelin ingestion appears to have subsided. Treatments at earlier than 4 months of infection have not promoted remyelination, suggesting that lesions reach a state of maturation, which poises them for repair.

10 However, without being limited to any specific mechanism, antibodies against glatiramer acetate appear to promote remyelination through an immunomodulatory mechanism, rather than through direct 15 stimulation of oligodendrocytes or their progenitors. It seems that antibodies against glatiramer acetate bind to activated microglia, macrophages, T cells and possibly to other similar types of cells, such as dendritic cells. Binding to these cells may modulate their functions, thereby 20 facilitating remyelination by oligodendrocytes. This hypothesis is supported by numerous findings of the subject invention. For example, in human glial cell cultures, antibodies against glatiramer acetate appeared to bind microglia by morphological criteria and by the criteria of 25 co-immunostaining with activated microglia markers. The glatiramer acetate-positive cells were distinct from O4-positive oligodendrocytes, which were extensively arborized. Glatiramer acetate antibodies also bound to perivascular infiltrates, further supporting the theory that glatiramer 30 acetate antibodies influence the activity of microglia or macrophages. The binding of antibodies against glatiramer acetate to glial cells differs from other remyelination-promoting antibodies, which bind to oligodendrocytes. Potential effects of the binding of antibodies against 35 glatiramer acetate include alteration in antigen

presentation, lymphocyte proliferation, and cytokine/growth factor production. In further support of this hypothesis, antibodies against glatiramer acetate were found to stimulate lymphocyte proliferation in vitro.

5

Most remyelination-promoting antibodies, including SCH 94.03 (1) are polyreactive autoantibodies derived from germline sequences (11, 42). Therefore, another intriguing possibility is that the complex

10 mixture of peptides comprising glatiramer acetate generated antibodies that mimicked protective, "natural" autoantibodies. In addition, antibodies against glatiramer acetate also had very low polyreactivity to other proteins, unlike other remyelination-promoting antibodies (e.g., SCH 15 94.03 monoclonal antibody), which cross-react with many protein antigens (1). Furthermore, antibodies against glatiramer acetate comprised a spectrum of antibody isotypes, unlike the predominance of IgMs in other remyelination-promoting antibodies (1). Another possibility is that 20 glatiramer acetate antibodies help to clear debris from lesions through opsonization, thereby permitting spontaneous remyelination to occur more readily.

An apparent paradox is that adoptive transfer of glatiramer 25 acetate antibodies promoted remyelination, yet active immunization did not. In fact, high-dose immunization increased lesion load. This suggests that glatiramer acetate had multiple effects in vivo and that the positive influence of the antibodies was overridden by other effects 30 of active immunization. The most commonly documented effect of glatiramer acetate is suppression of Th1 lymphocyte activity through mechanisms such as induction of Th2 suppressor lymphocytes, inhibition of peptide binding to MHC Class II, and T cell antagonism (8-10, 18, 21, 51, 67). 35 Since glatiramer acetate-mediated suppression is relatively

nonspecific (12, 20, 67, 76), and since T lymphocytes are essential for controlling Theiler's virus even during late disease (44, 61), it is possible that antiviral immunity was depressed by glatiramer acetate, resulting in increased viral 5 pathogenesis and lesion exacerbation. This hypothesis is consistent with the increased virus antigen expression, decreased antiviral antibody titers, and decreased TMEV-specific DTH responses. These changes were mild, raising the possibility that glatiramer acetate immunization also 10 expanded lesions through other mechanisms. If synergy between glatiramer acetate antibodies and macrophages or lymphocytes is required for remyelination, then an alteration in the function of these cells by glatiramer acetate (34, 49) could have abrogated remyelination.

15

A dogma in the multiple sclerosis field is that immune activation, both cellular and humoral, exerts an overwhelmingly deleterious role and must be suppressed for effective therapy. However, it is becoming increasingly clear 20 that the immune system can also be protective in the injured CNS through mechanisms such as secretion of trophic factors (25, 29, 43, 54). For example, following experimental optic nerve damage, glatiramer acetate reactive and MBP reactive lymphocytes reduce secondary neuronal degeneration (43). 25 Myelin repair by glatiramer acetate antibodies exemplifies the therapeutic contribution that humoral activation can make. Maximizing the humoral response to glatiramer acetate through various means can enhance the restoration of conduction and axon health after acute demyelinating attacks.

30

REFERENCES

1. U.S. Patent No. 5,591,629, issued January 7, 1997 (Rodriguez et al.).
- 5 2. U.S. Patent No. 5,800,808, issued September 1, 1998 (Konfino et al.).
- 10 3. U.S. Patent No. 5,981,589, issued November 9, 1999 (Konfino et al.).
- 15 4. U.S. Patent No. 6,048,898, issued April 11, 2000 (Konfino et al.).
5. U.S. Patent No. 6,054,430, issued April 25, 2000 (Konfino et al.).
- 15 6. WO 00/05250, published February 3, 2000 (Aharoni et al.).
- 20 7. Aharoni, R. et al. 2000. Specific Th2 cells accumulate in the central nervous system of mice protected against experimental autoimmune encephalomyelitis by copolymer 1. Proc. Natl. Acad. Sci. USA 97: 11472-11477.
- 25 8. Aharoni, R., D. Teitelbaum, M. Sela and R. Arnon. 1997. Copolymer 1 induces T cells of the T helper type 2 that crossreact with myelin basic protein and suppress experimental autoimmune encephalomyelitis. Proc. Natl. Acad. Sci. USA 94: 10821-10826.
- 30 9. Aharoni, R., D. Teitelbaum and R. Arnon. 1993. T suppressor hybridomas an interleukin-2-dependent lines induce by copolymer 1 or by spinal cord homogenate down-regulate experimental allergic encephalomyelitis. Eur. J. Immunol. 23:17-25.

10. Aharoni, R., D. Teitelbaum, R. Arnon and M. Sela. 1999. Copolymer 1 acts against the immunodominant epitope 82-100 of myelin basic protein by T cell receptor antagonism in addition to major histocompatibility complex blocking. Proc. 5 Natl. Acad. Sci. USA 96: 634-639.

11. Asakura, K., D.J. Muller, R.J. Pogulis, L.R. Pease and M. Rodriguez, 1995. Oligodendrocyte-reactive O1, O4, and HNK-1 monoclonal antibodies are encoded by germline 10 immunoglobulin genes. Molec. Brain Res. 34: 283-293.

12. Ben-Nun, A. et al. 1996. The autoimmune reactivity to myelin oligodendrocyte glycoprotein (MOG) in multiple sclerosis is potentially pathogenic: effect of Copolymer 1 on 15 MOG-induced disease. J. Neurol. 243 (Suppl 1): S14-S22.

13. Bielekova, B. et al. 2000. Encephalitogenic potential of the myelin basic protein peptide amino acids 83-99. in multiple sclerosis: Results of a phase II clinical trial with 20 an altered peptide ligand. Nat. Med. 6: 1167-1175.

14. Compston, D.A.S. 1991. "Genetic susceptibility to multiple sclerosis," in McAlpine's Multiple Sclerosis, Matthews, B. ed., London: Churchill Livingstone, 301-319. 25

15. Dal Canto, M.C., and H.L. Lipton. 1977. Multiple sclerosis. Animal model: Theiler's virus infection in mice. Am. J. Path. 88:497-500.

30 16. Darnell et al. 1990. Molecular Cell Biology, 2nd ed., New York: Scientific American Books, 172, 768, 778-79, 1038, 1040.

35 17. Duda, P. W., M.C. Schmied, S.L. Cook, J.I. Krieger and D.A. Hafler, 2000. Glatiramer acetate Copaxone®. induces

degenerate, Th2-polarized immune responses in patients with multiple sclerosis. *J. Clin. Invest.* 105: 967-976.

18. Fridkis-Hareli, M. et al. 1994. Direct binding of myelin basic protein and synthetic copolymer 1 to class II major histocompatibility complex molecules on living antigen-presenting cells -- specificity and promiscuity. *Proc. Natl. Acad. Sci. USA* 91: 4872-4876.

10 19. Fridkis-Hareli, M., D. Teitelbaum, E. Gurevich, I. Pecht, C. Brautbar, O.J. Kwon, T. Brenner, R. Arnon, and M. Sela. 1994. Direct binding of myelin basic protein and synthetic copolymer 1 to class II major histocompatibility complex molecules on living antigen-presenting cells-- specificity and promiscuity. *Proc. Natl. Acad. Sci. USA* 91:4872-4826.

20 20. Fridkis-Hareli, M., E.F. Rosloniec, L. Fugger, and J.L. Strominger. 1998. Synthetic amino acid copolymers that bind to HLA-DR proteins and inhibit type II collagen-reactive T cells clones. *Proc. Natl. Acad. Sci. USA* 95: 12528-12531.

25 21. Fridkis-Hareli, M. and J.L. Strominger, 1998. Promiscuous binding of synthetic copolymer 1 to purified HLA-DR molecules. *J. Immunol.* 160: 4386-4397.

22. Genain, C.P. et al. 1995. Antibody facilitation of multiple sclerosis-like lesions in a nonhuman primate. *J. Clin. Invest.* 96: 2966-2974.

30 23. Hafler, D.A. and H.L. Weiner. 1989. MS: A CNS and systemic autoimmune disease. *Immunol. Today* 10:104-107.

35 24. Hawes, G. et al. 1995. Limited restriction in the TCR-alpha beta V region usage of antigen-specific clones.

Recognition of myelin basic protein (amino acids 84-102) and *Mycobacterium bovis* 65-kDa heat shock protein (amino acids 3-13) by T cell clones established from peripheral blood mononuclear cells of monozygotic twins and HLA-identical 5 individuals. *J. Immunol.* 154:2, 555-566.

25. Huang, D.W., L. McKerracher, P.E. Braun, and S. David, 1999. A therapeutic vaccine approach to stimulate axon regeneration in the adult mammalian spinal cord. *Neuron* 24: 10 639-647.

26. Johnson, K.P., B.R. Brooks, J.A. Cohen, C.C. Ford, J. Goldstein, R.P. Lisak, L.W. Myers, H.S. Panitch, J.W. Rose, R.B. Schiffer et al. 1995. Copolymer 1 reduces relapse rate 15 and improves disability in relapsing-remitting multiple sclerosis: results of a phase III multicenter, double-blind placebo-controlled trial. The Copolymer 1 Multiple Sclerosis Study Group. *Neurol.* 45:1268.

20 27. Johnson, K.P. et al. 2000. Sustained clinical benefits of glatiramer acetate in relapsing multiple sclerosis patients observed for 6 years. Copolymer 1 Multiple Sclerosis Study Group. *Mult. Scler.* 6: 255-266.

25 28. Kappos, L. et al. 2000. Induction of a non-encephalitogenic type 2 T helper-cell autoimmune response in multiple sclerosis after administration of an altered peptide ligand in a placebo-controlled, randomized phase II trial. *Nat. Med.* 6: 1176-1182.

30 29. Kerschensteiner, M. et al. 1999. Activated human T cells, B cells, and monocytes produce brain-derived neurotrophic factor in vitro and in inflammatory brain lesions: a neuroprotective role of inflammation?: *J. Exp. Med.* 189: 865-870.

30. Kipnis, J. et al. 2000. T cell immunity to copolymer 1 confers neuroprotection on the damaged optic nerve: possible therapy for optic neuropathies. Proc. Natl. Acad. Sci. USA 97: 7446-7451.

5

31. Lampert, P.W. 1978. Autoimmune and virus-induced demyelinating diseases. A review. Am. J. Path. 91:176-208.

10

32. Lando, Z., D. Teitelbaum and R. Arnon. 1979. Effect of cyclophosphamide on suppressor cell activity in mice unresponsive to EAE. J. Immunol. 123: 2156-2160.

15

33. Lehrich, J. and B. Arnason. 1976. Demyelinative myelopathy in mice induced by the DA virus. J. Neurol Sci. 29:149.

20

34. Li, Q., R. Milo, H. Panitch,, P. Swoveland, and C.T. Bever, Jr. 1998. Glatiramer acetate blocks the activation of THP-1 cells by interferon-gamma. Eur. J. Pharmacol. 342: 303-310.

25

35. Linington, C., M. Bradl,, H. Lassmann, C. Brunner, and K. Vass. 1988. Augmentation of demyelination in rat acute allergic encephalomyelitis by circulating mouse monoclonal antibodies directed against a myelin/oligodendrocyte glycoprotein. Am. J. Pathol. 130: 443-454.

30

36. Lipton, H.L. and M.C. Dal Canto. 1976. Chronic neurologic disease in Theiler's virus infection of SJL/J mice. J. Neurol. Sci. 30: 201-207.

35

37. Lisak, R.P. et al. 1983. Effect of treatment with Copolymer 1 (Cop-1) on the in vivo and in vitro manifestations of Experimental Allergic Encephalomyelitis (EAE). J. Neurol. Sci. 62: 281-293.

38. Martyn, C. 1991. "The epidemiology of multiple sclerosis" in McAlpine's Multiple Sclerosis, Matthews, B., ed., London: Churchill Livingstone, 3-40.

5 39. McGavern, D.B. et al. 1999. Quantitation of spinal cord demyelination, remyelination, atrophy, and axonal loss in a model of progressive neurologic injury. *J. Neurosci. Res.* 58: 492-504.

10 40. Miller, D.J., K. Asakura, and M. Rodriguez. 1995. Experimental strategies to promote central nervous system remyelination in multiple sclerosis: insights gained from the Theiler's virus model system. [Review]. *J. Neurosci. Res.* 41:291.

15 41. Miller, D.J., K. S. Sanborn, J. A. Katzmann, and M. Rodriguez. 1994. Monoclonal autoantibodies promote central nervous system repair in an animal model of multiple sclerosis. *J. Neurosci.* 14:6230-6238.

20 42. Miller, D.J. and M. Rodriguez. 1995. A monoclonal autoantibody that promotes central nervous system remyelination in a model of multiple sclerosis is a natural autoantibody encoded by germline immunoglobulin genes. *J. Immunol.* 154:2460-2469.

25 43. Moalem, G. et al. 1999. Autoimmune T cells protect neurons from secondary degeneration after central nervous system axotomy. *Nat. Med.* 5: 49-55.

30 44. Murray, P.D., K.D. Pavelko, J. Leibowitz, X. Lin, and M. Rodriguez. 1998. CD4+ and CD8+ T cells make discrete contributions to demyelination and neurologic disease in a viral model of multiple sclerosis. *J. Virol.* 72: 7320-7329.

35

45. Neuhaus, O. et al. 2000. Multiple sclerosis: comparison of copolymer-1-reactive T cell lines from treated and untreated subjects reveals cytokine shift from T helper 1 to T helper 2 cells. *Proc. Natl. Acad. Sci. USA* 97: 7452-7457.

5

46. Neville, K.L., M.C. Dal Canto, J.A. Bluestone and S.D. Miller. 2000. CD28 costimulatory blockade exacerbates disease severity and accelerates epitope spreading in a virus-induced autoimmune disease. *J. Virol.* 74: 8349-8357.

10

47. Olsson, T. 1992. Immunology of multiple sclerosis. *Curr. Opin. Neurol. Neurosurg.* 5:195-202.

48. Pavelko, K.D., B.G. van Engelen and M. Rodriguez. 1998. Acceleration in the rate of CNS remyelination in lyssolecithin-induced demyelination. *J. Neurosci.* 18: 2498-2505.

15

49. Pratt, A., A. Al Asmi, P. Duquette and J.P. Antel. 1999. Lymphocyte migration and multiple sclerosis: relation with disease course and therapy. *Ann. Neurol.* 46: 253-256.

20

50. Qin, Y., D.Q. Zhang, A. Prat, S. Pouly, and J. Antel. 2000. Characterization of T cell lines derived from glatiramer-acetate-treated multiple sclerosis patients. *J. Neuroimmunol.* 108: 201-206.

25

51. Racke, M.K., R. Martin, H. McFarland, and R.B. Fritz. 1992. Copolymer-1-induced inhibition of antigen-specific T cell activation: interference with antigen presentation. *J. Neuroimmunol.* 37:75.

30

52. Rader, C. and C. Barbas. 1997. Phage display of combinatorial antibody libraries. *Curr. Opin. Biotech.* 8:503-508.

35

53. Raine, C.S., Cannella, B., Hauser, S.L. and Genain, C.P. 1999. Demyelination in primate autoimmune encephalomyelitis and acute multiple sclerosis lesions: a case for antigen-specific antibody mediation [see comments]. *Ann. Neurol.* 46: 144-146.

5 54. Rapalino, O. et al. 1998. Implantation of stimulated homologous macrophages results in partial recovery of paraplegic rats. *Nat. Med.* 4: 814-821.

10 55. Rodriguez, M. 1991. Immunoglobulins stimulate central nervous system remyelination: electron microscopic and morphometric analysis of proliferating cells. *Lab. Invest.* 64:358.

15 56. Rodriguez, M. 1992. Central nervous system demyelination and remyelination in multiple sclerosis and viral models of disease. *J. Neuroimmunol.* 40:255.

20 57. Rodriguez, M. and B. Scheithauer. 1994. Ultrastructure of multiple sclerosis. *Ultrastruct. Pathol.* 18:3.

25 58. Rodriguez, M., D.J. Miller, and V.A. Lennon. 1996. Immunoglobulins reactive with myelin basic protein promote CNS remyelination. *Neurol.* 46:538-545.

30 59. Rodriguez, M. et al. 1987. Theiler's murine encephalomyelitis: a model of demyelination and persistence of virus. *Crit. Rev. Immunol.* 7:325.

60. Rodriguez, M., L.M. Siegel, D. Hovanec-Burns, L. Bologa and M.C. Graves. 1988. Theiler's virus-associated antigens on the surfaces of cultured glial cells. *Virol.* 166: 463-474.

35 61. Rodriguez, M. and M.D. Lindsley. 1992. Immunosuppression

promotes CNS remyelination in chronic virus-induced demyelinating disease. *Neurol.* 42: 348-357.

62. Rodriguez, M. and V.A. Lennon. 1990. Immunoglobulins
5 promote remyelination in the central nervous system. *Ann.*
Neurol. 27:12.

63. Rodriguez, M., V.A. Lennon, E.N. Benveniste, and J.E.
Merrill. 1987. Remyelination by oligodendrocytes stimulated
10 by antiserum to spinal cord. *J. Neuropathol. Exp. Neurol.*
46:84.

64. Sela, M. et al. 1990. Suppressive activity of Cop-1 in
EAE and its relevance to multiple sclerosis. *Bull. Inst.*
15 *Pasteur.* 88:303-314.

65. Teitelbaum, D., A. Meshorer, T. Hirshfeld, R. Arnon and
M. Sela. 1971. Suppression of experimental allergic
encephalomyelitis by a synthetic polypeptide. *Eur. J.*
20 *Immunol.* 1: 242-248.

66. Teitelbaum, D., C. Webb, A. Meshorer, R. Arnon and M.
Sela. 1973. Suppression by several synthetic polypeptides of
experimental allergic encephalomyelitis induced in guinea
25 pigs and rabbits with bovine and human basic encephalitogen.
Eur. J. Immunol. 3: 273-279.

67. Teitelbaum, D. et al. 1996. Copolymer 1 inhibits chronic
relapsing experimental allergic encephalomyelitis induced by
30 proteolipid protein (PLP) peptides in mice and interferes
with PLP-specific T cell responses. *J. Neuroimmunol.* 64: 209-
217.

68. Teitelbaum, D., R. Aharoni, M. Sela and R. Arnon. 1991.
35 Cross-reactions and specificities of monoclonal antibodies

against myelin basic protein and against the synthetic Copolymer 1. Proc. Natl. Acad. Sci. USA 88:9528-9532.

5 69. Teitelbaum, D., R. Milo, R. Arnon and M. Sela. 1992. Synthetic copolymer 1 inhibits human T cell lines specific for myelin basic protein. Proc. Natl. Acad. Sci. USA 89:137.

10 70. Tuohy, V.K. et al. 1989. Identification of an encephalitogenic determinant of myelin proteolipid protein for SJL mice. J. Immunol. 142:5, 1523-1527.

15 71. Van Engelen, B.G., D.J. Miller, K.D. Pavelko, O.R. Hommes, and M. Rodriguez. 1994. Promotion of remyelination by polyclonal immunoglobulin in Theiler's virus-induced demyelination and in multiple sclerosis. [Review]. J. Neurol. Neurosurg. Psych. 57(Suppl):65.

20 72. Vaswani, S.K. et al. 1998. Humanized antibodies as potential therapeutic drugs. Ann. Allergy Asthma Immunol. 81:105-119.

25 73. Warrington, A.E. et al. 2000. Human monoclonal antibodies reactive to oligodendrocytes promote remyelination in a model of multiple sclerosis. Proc. Natl. Acad. Sci. USA 97: 6820-6825.

30 74. Webb, C. et al. 1973. In vivo and in vitro immunological cross-reactions between basic encephalitogen and synthetic basic polypeptides capable of suppressing Experimental Allergic Encephalomyelitis. Eur. J. Immunol. 3: 279-286.

75. Webb, C. et al. 1976. Molecular requirements involved in suppression of EAE by synthetic basic copolymers of amino acids. Immunochem. 13:333-337.

76. Zhang, M. et al. 2000. Copolymer 1 inhibits experimental autoimmune uveoretinitis. *J. Neuroimmunol.* 103: 189-194.

77. "Copaxone" in Physician's Desk Reference, 2000, Medical
5 Economics Co., Inc., Montvale, NJ, 3115.

78. Production of monoclonal antibodies. 1991. *Curr. Protocols.* Unit 2.5.1 - 2.5.17.